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Zukowski

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(54) **BRANCHED STENT DELIVERY SYSTEM**

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A61F 2/06 (2006.01)

(52) **U.S. Cl.** **623/1.11**

(58) **Field of Classification Search** None
See application file for complete search history.

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Primary Examiner—Todd E Manahan

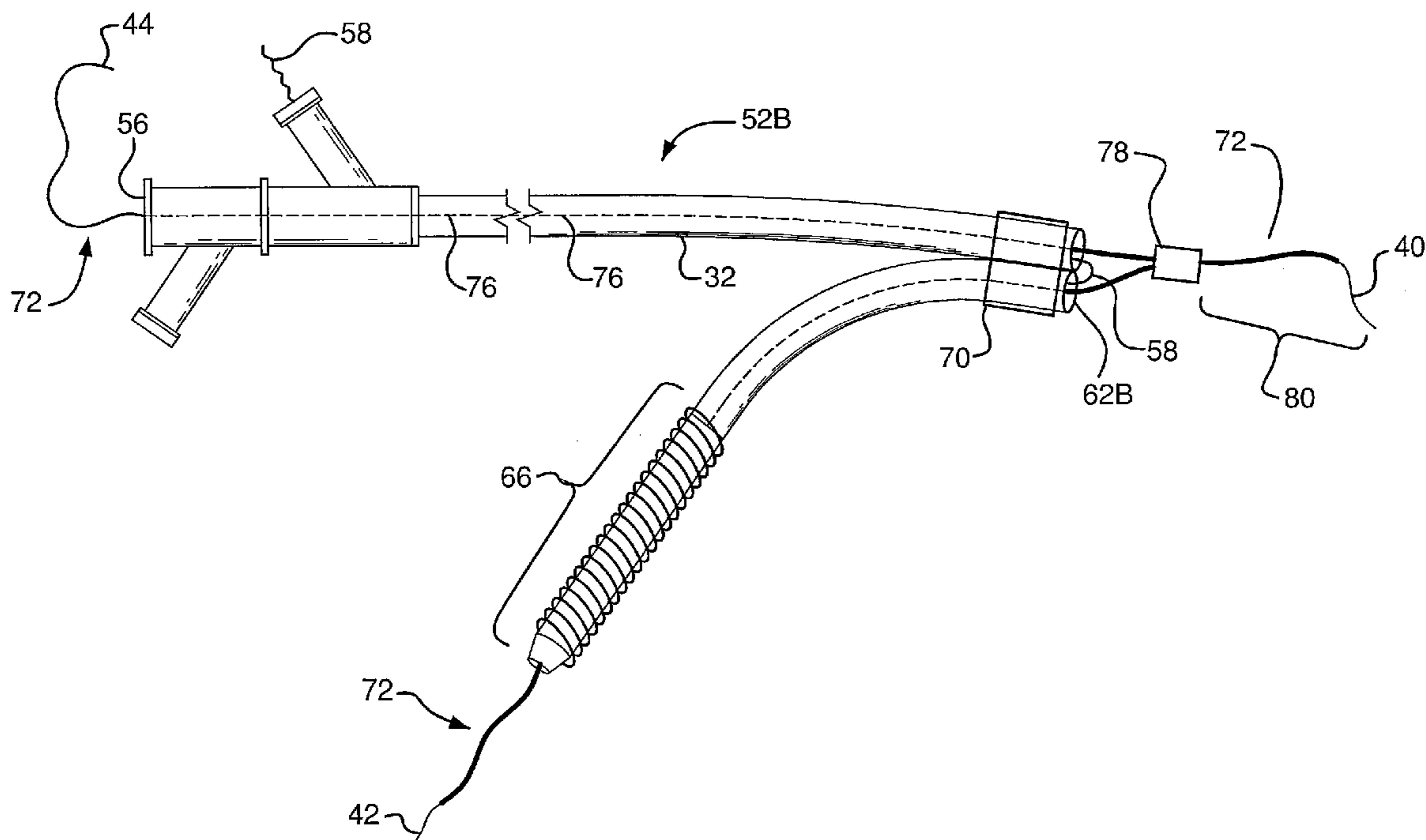
Assistant Examiner—Naquan Ishman

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(57) **ABSTRACT**

An interventional delivery system with a first catheter having at its distal end a side branch vessel segment; a second catheter attached around the first catheter and having at its distal end a main vessel segment; a side branch vessel device attached to side branch vessel segment of the first catheter; and main vessel device attached to the main vessel segment of the second catheter. The main vessel device and the side branch vessel device are able to be simultaneously delivered to a treatment site.

3 Claims, 18 Drawing Sheets



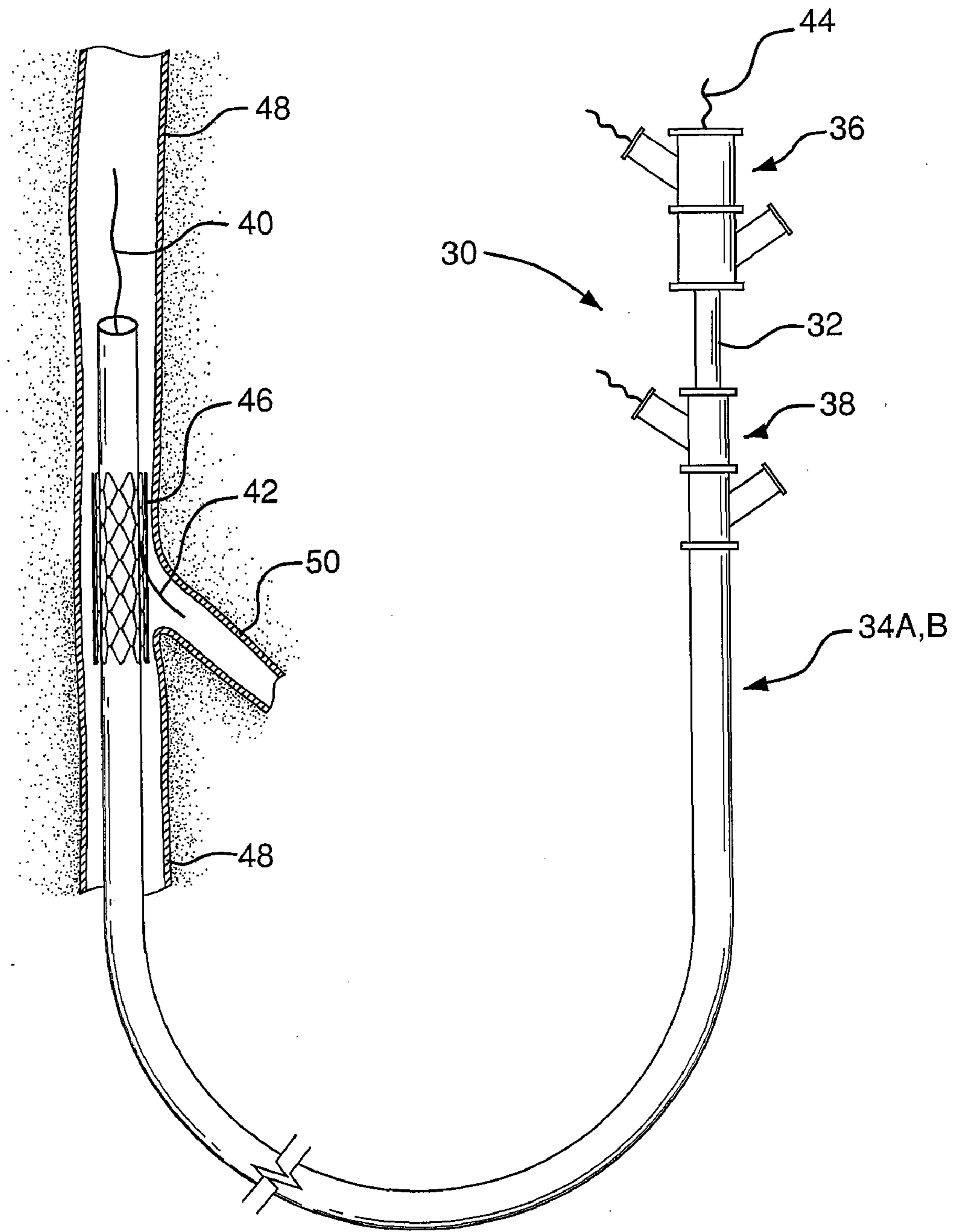


FIG. 1

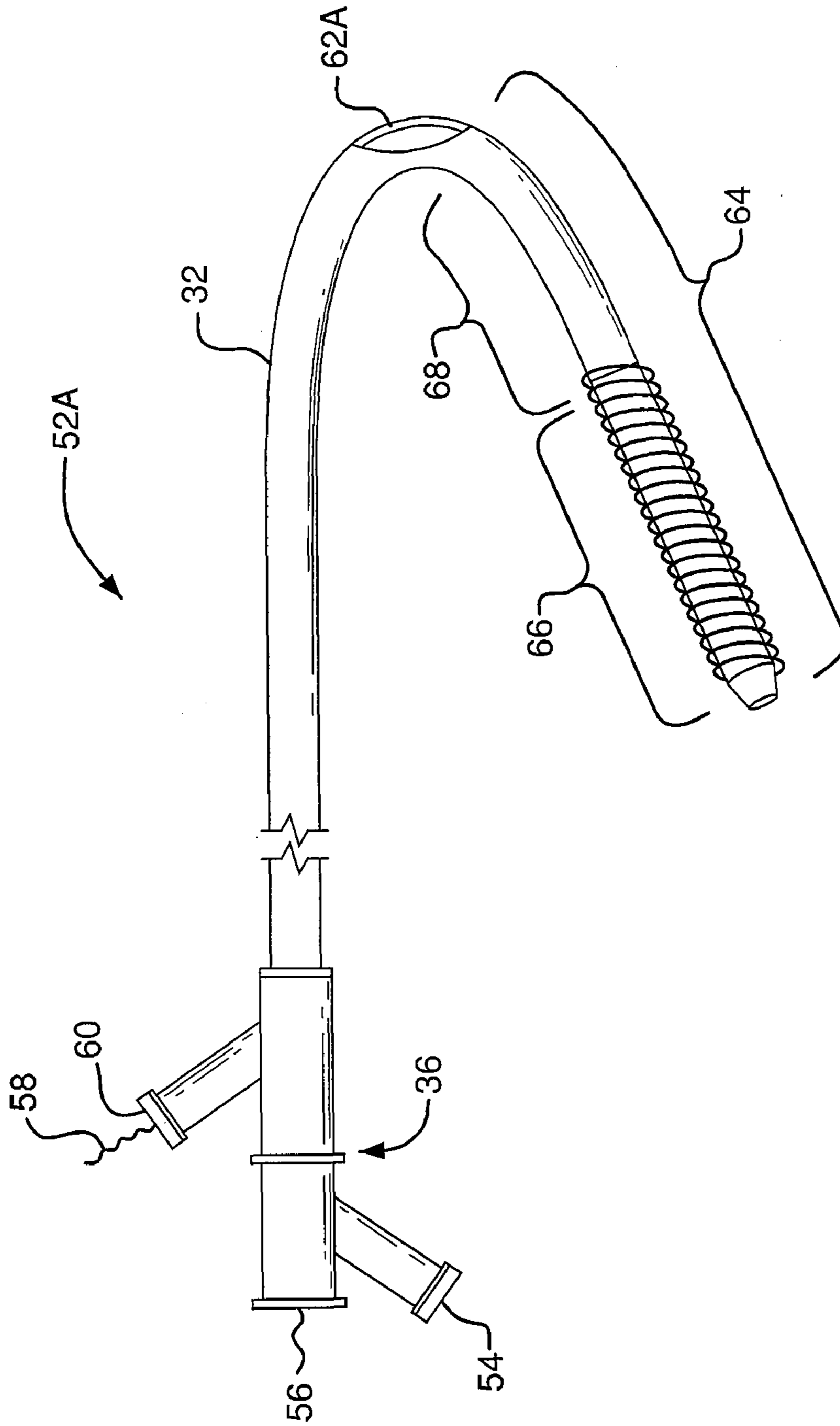


FIG. 2A

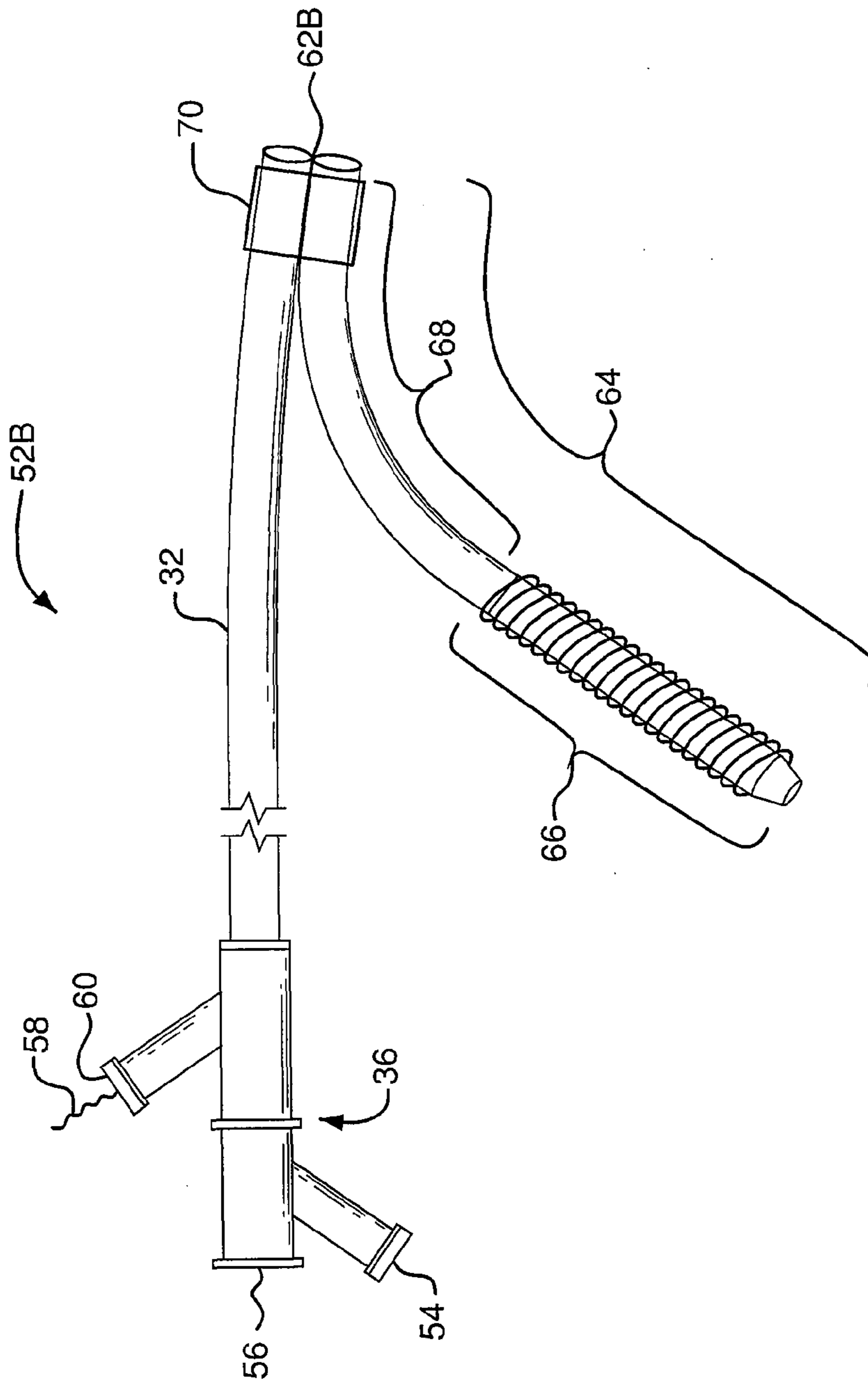


FIG. 2B

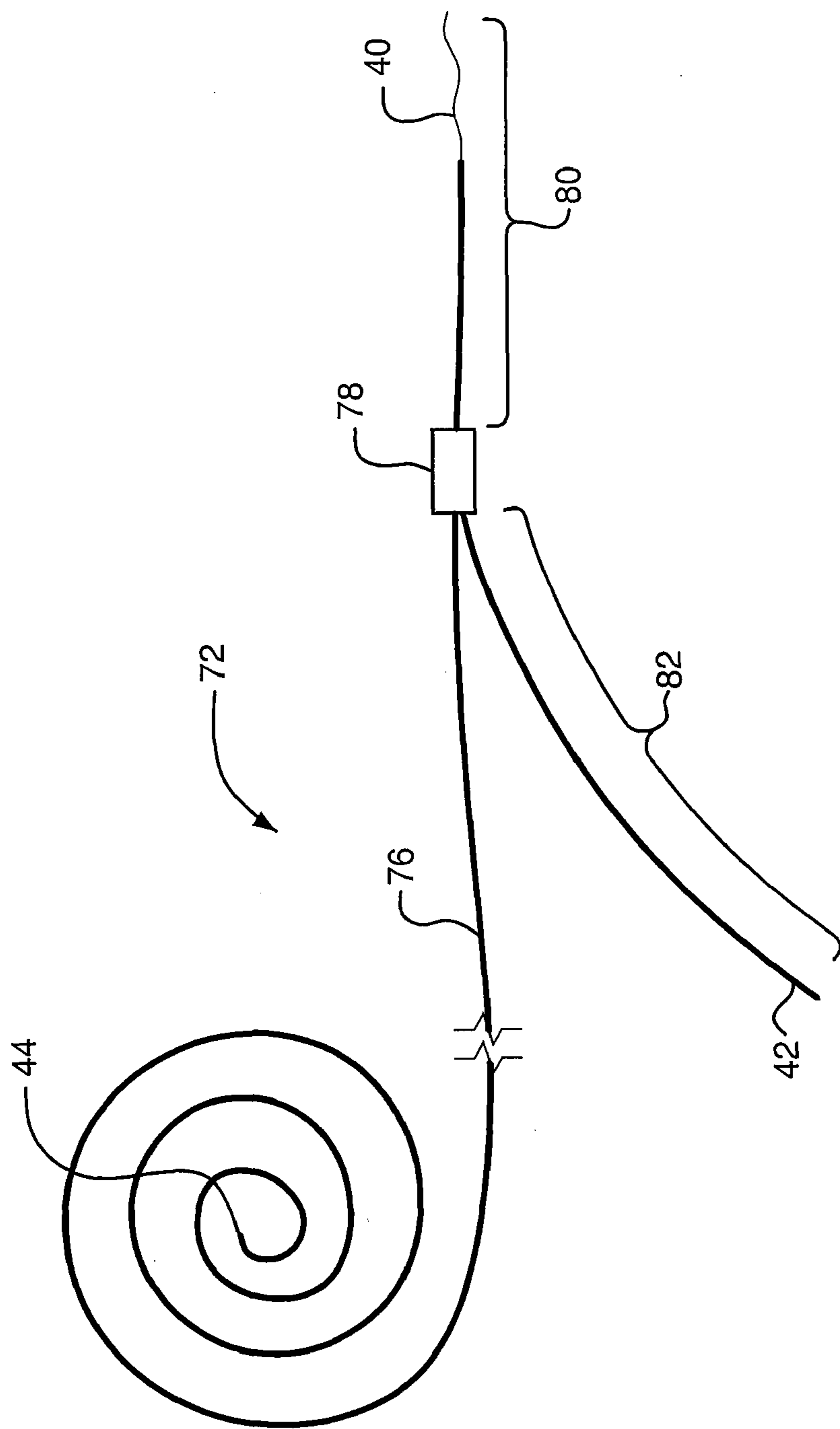


FIG. 3

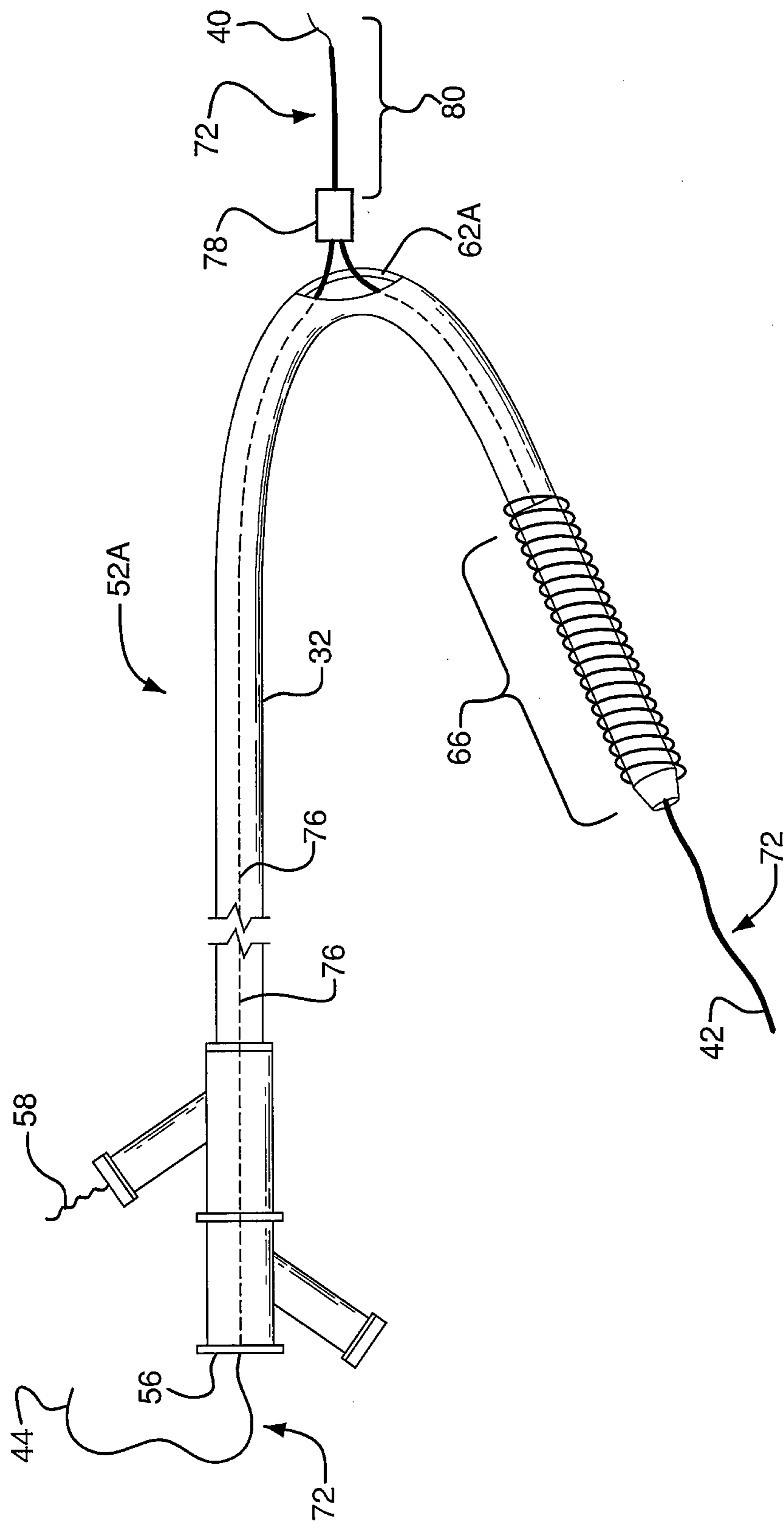


FIG. 4A

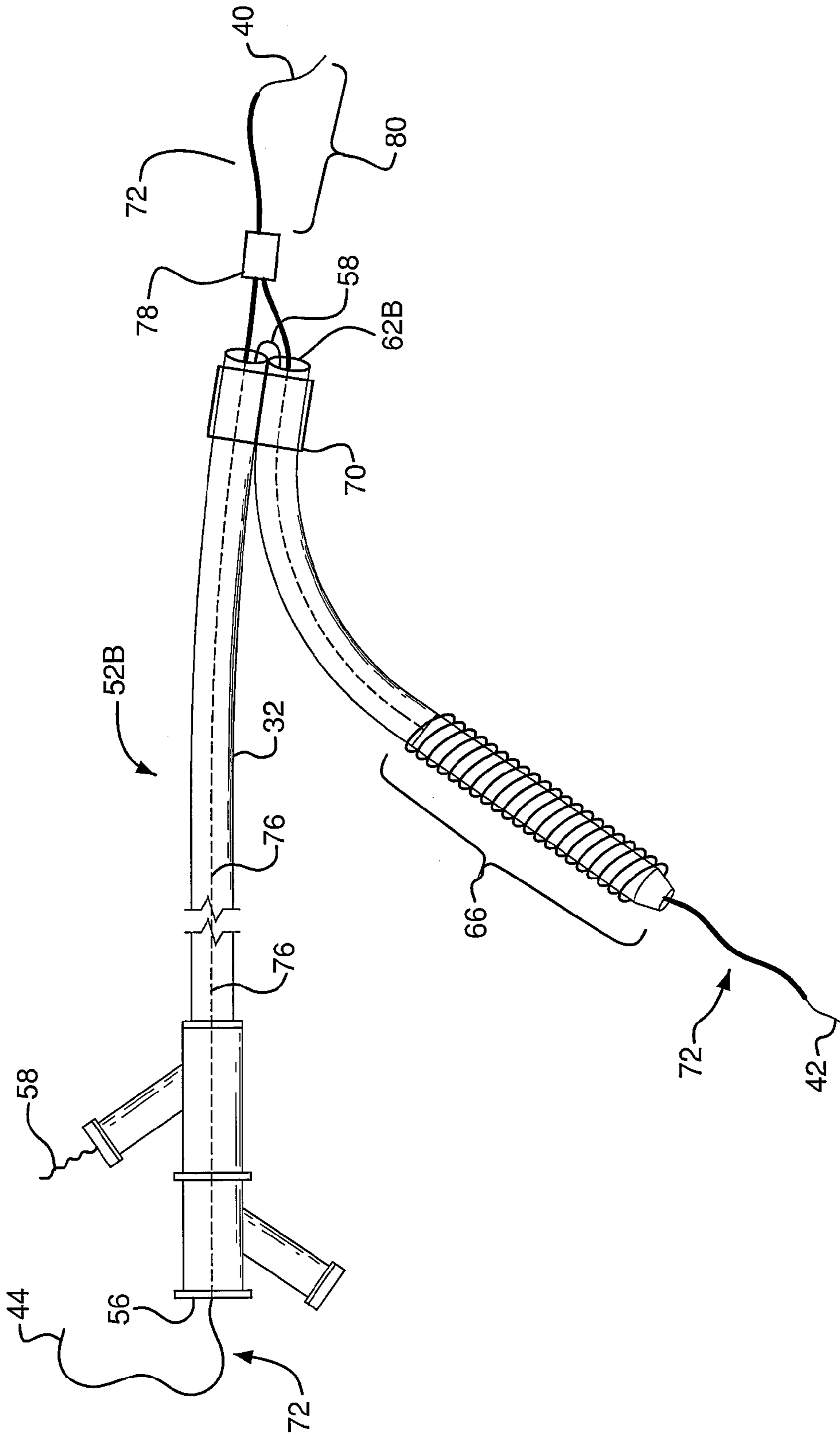


FIG. 4B

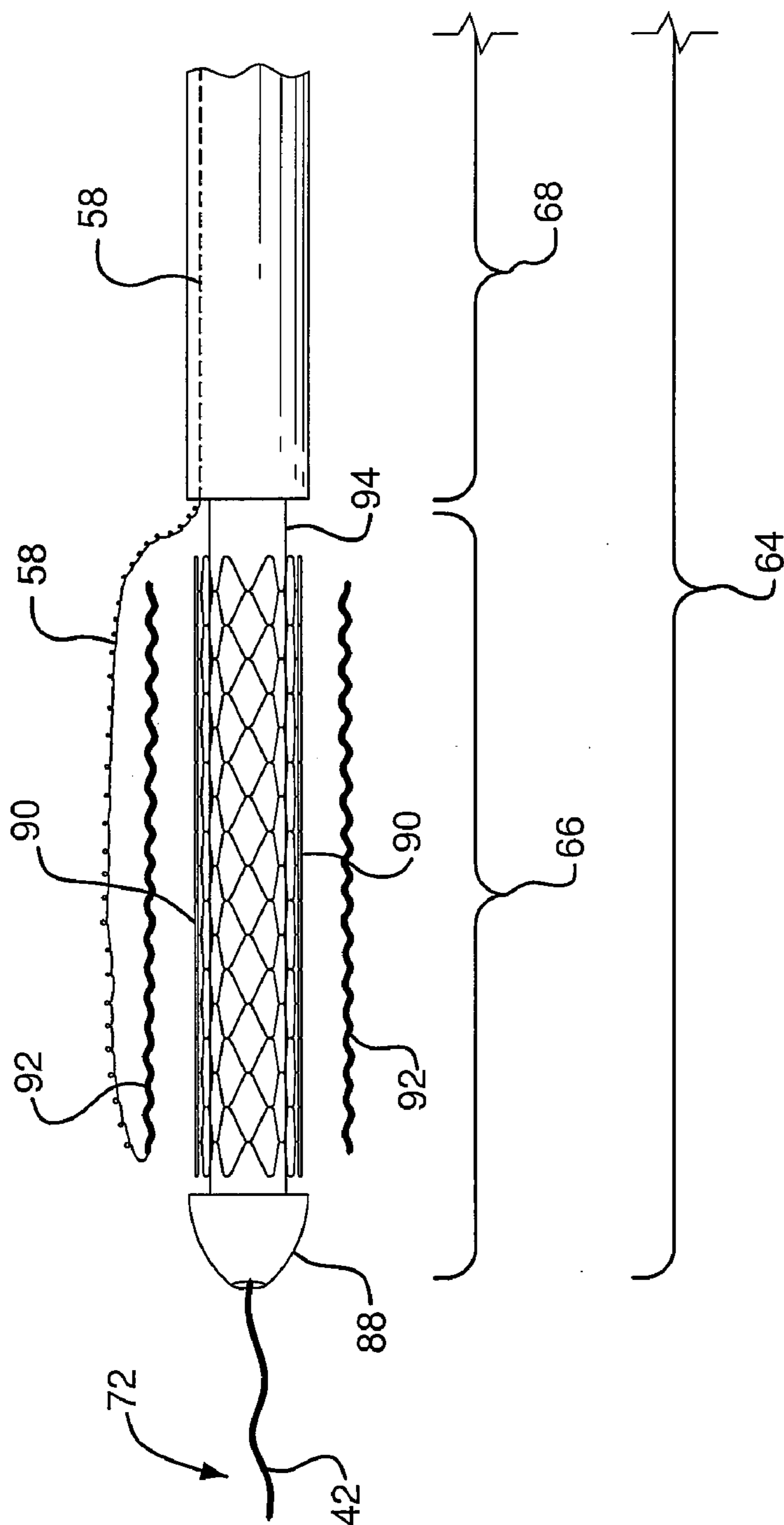


FIG. 5

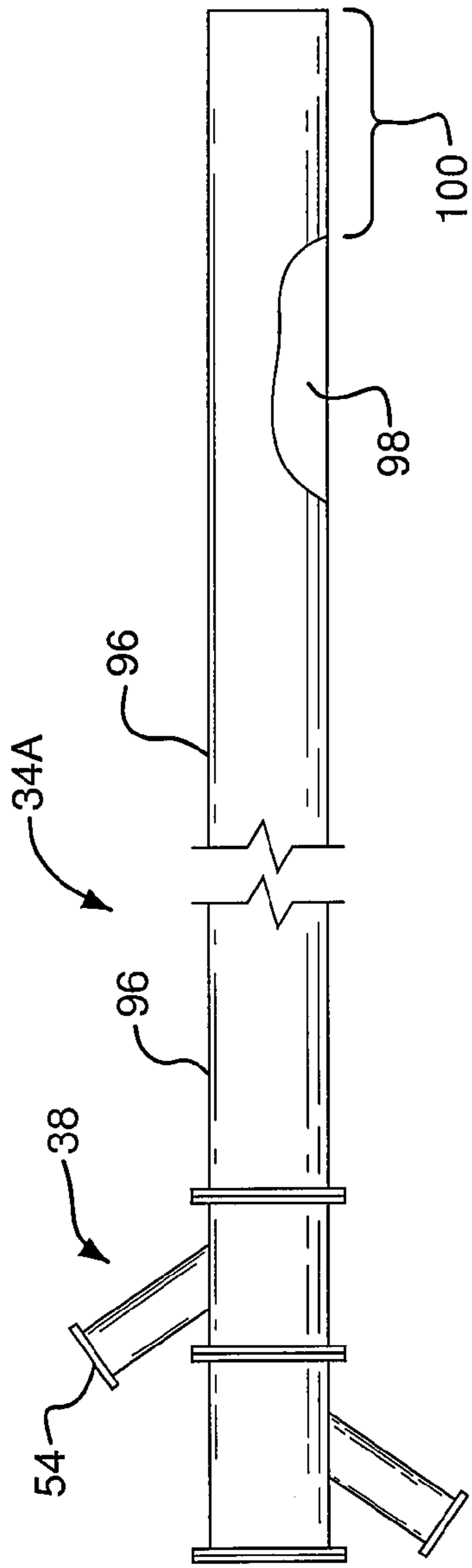


FIG. 6A

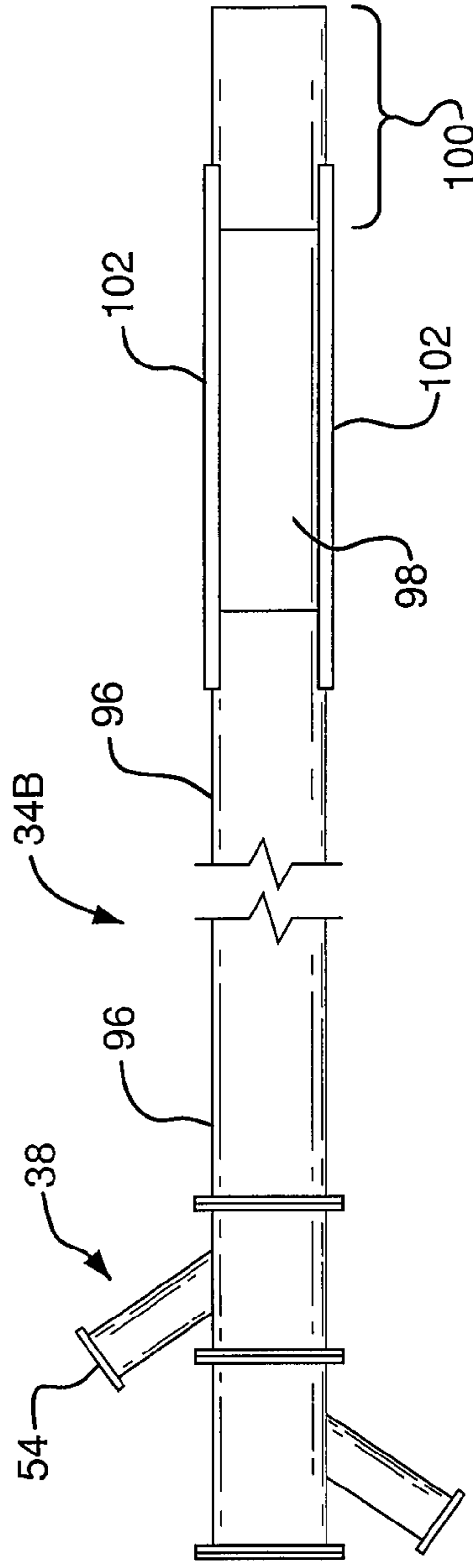


FIG. 6B

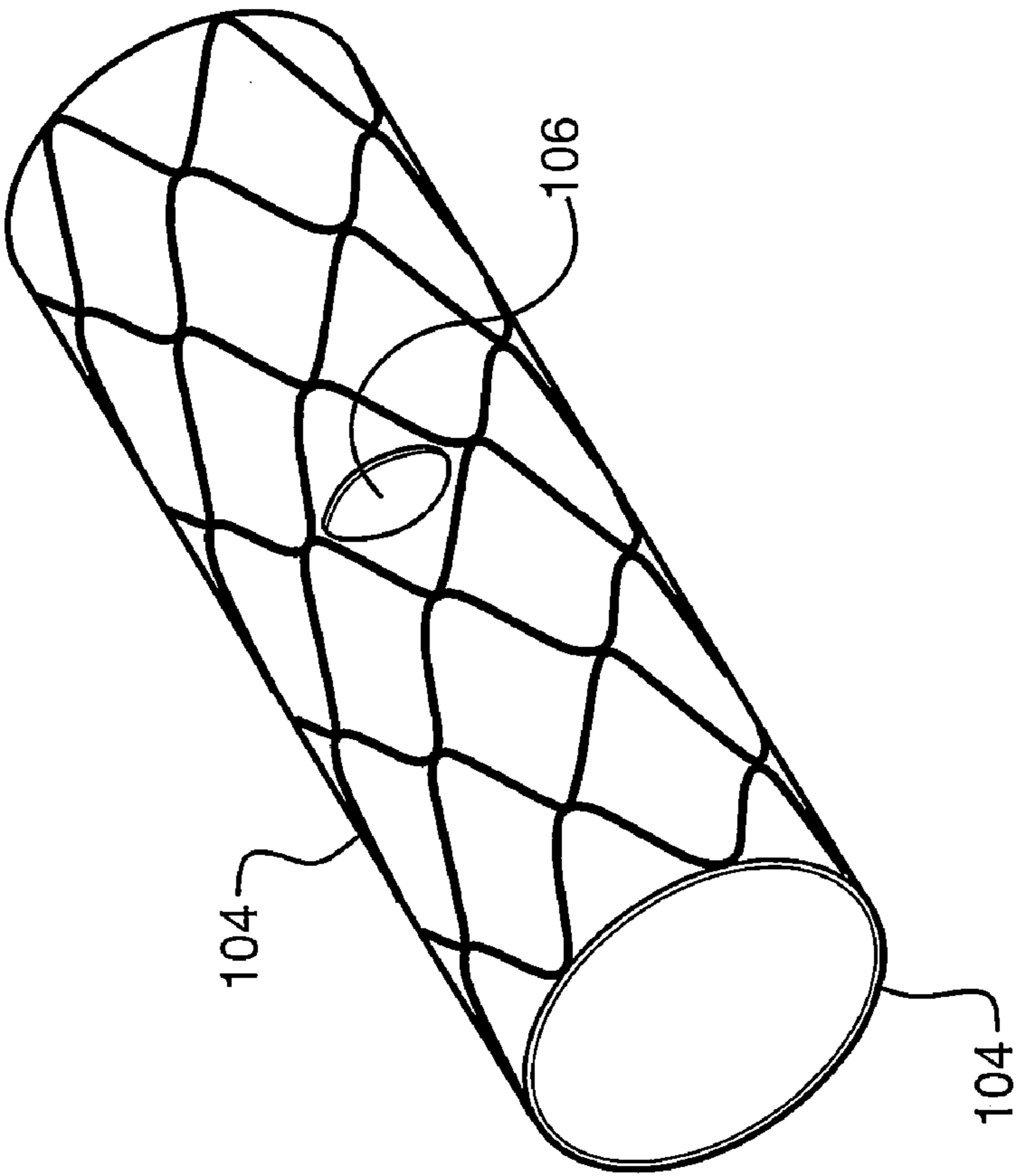


FIG. 7

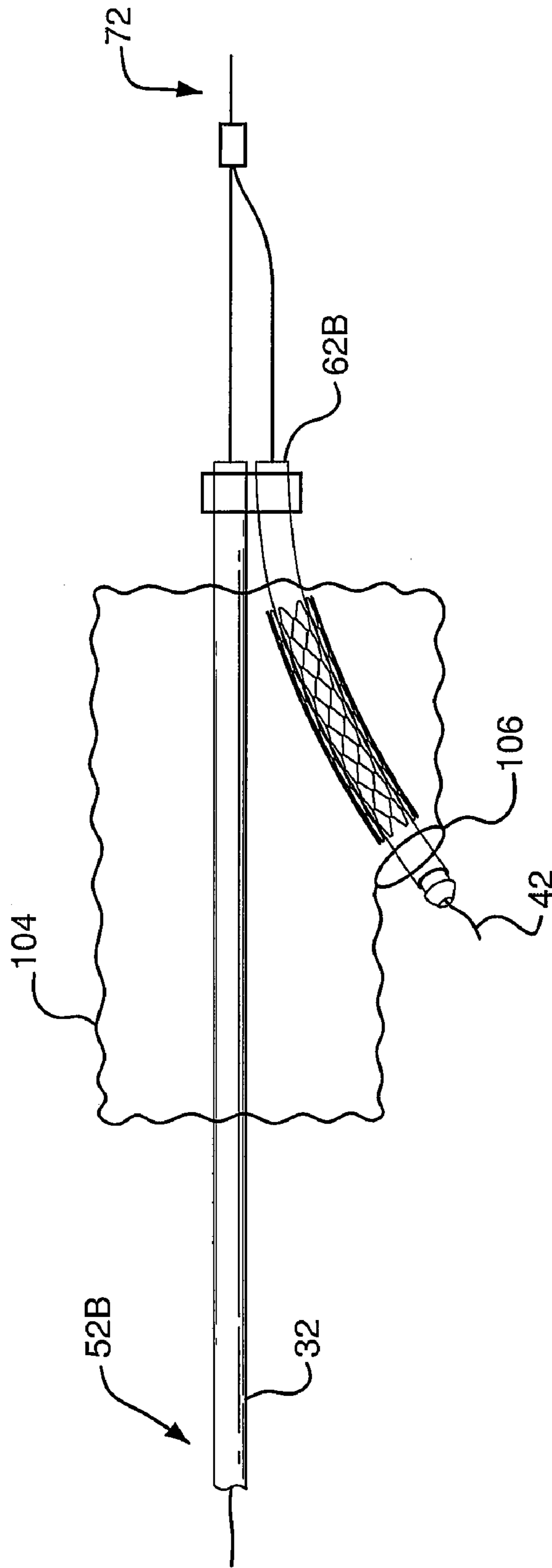


FIG. 8

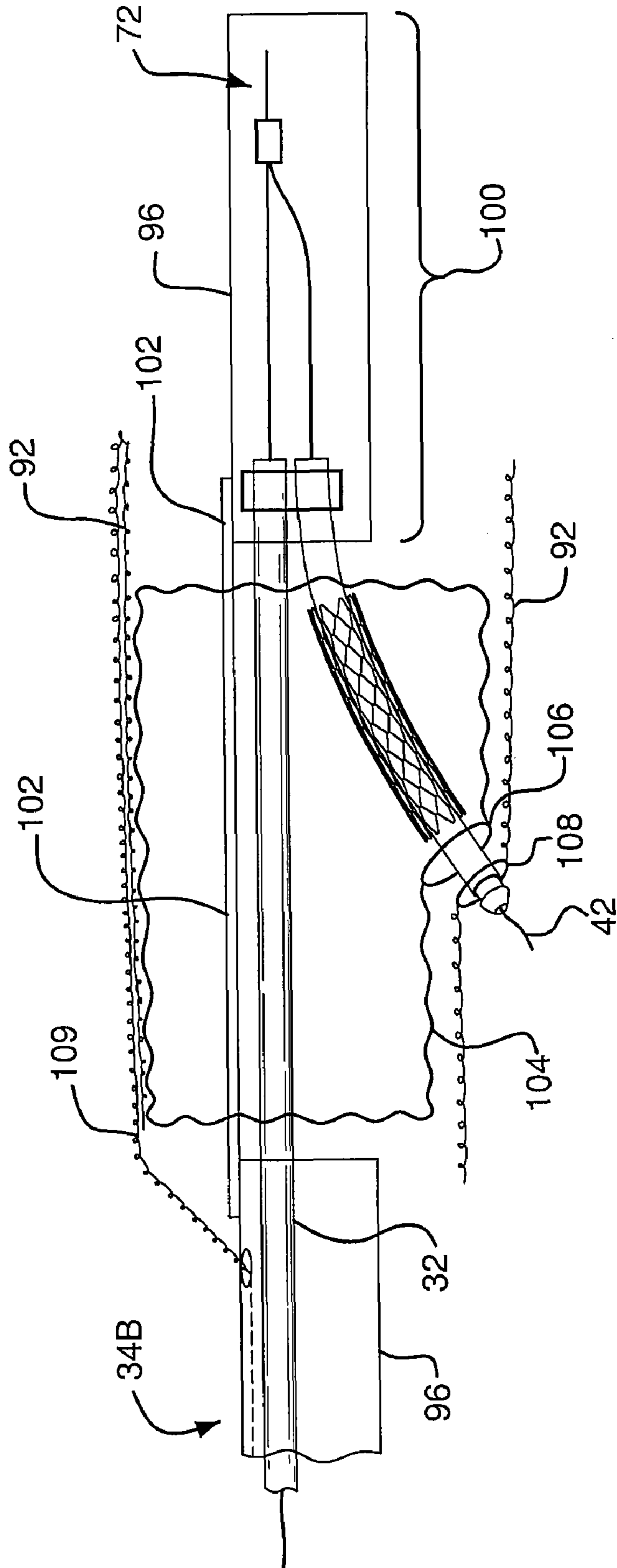


FIG. 9

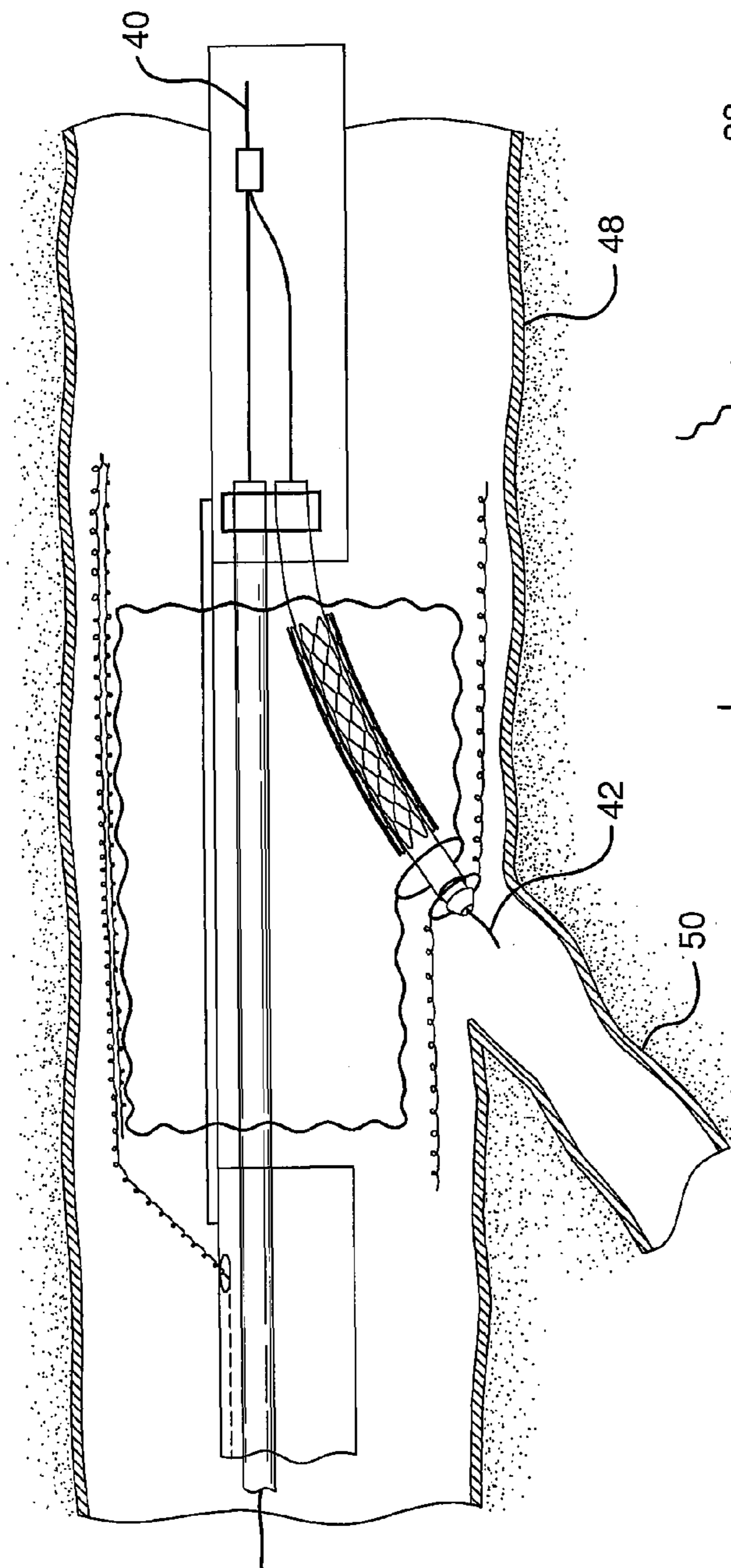


FIG. 10A

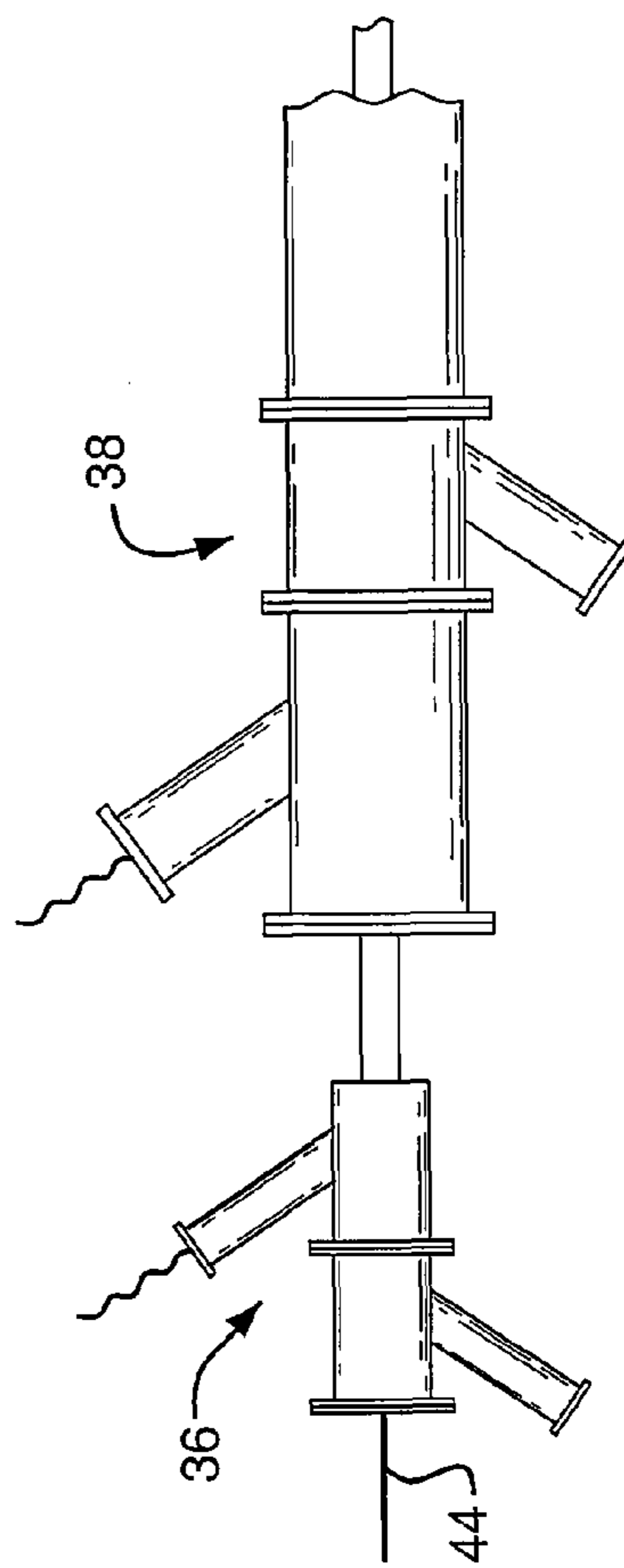


FIG. 10B

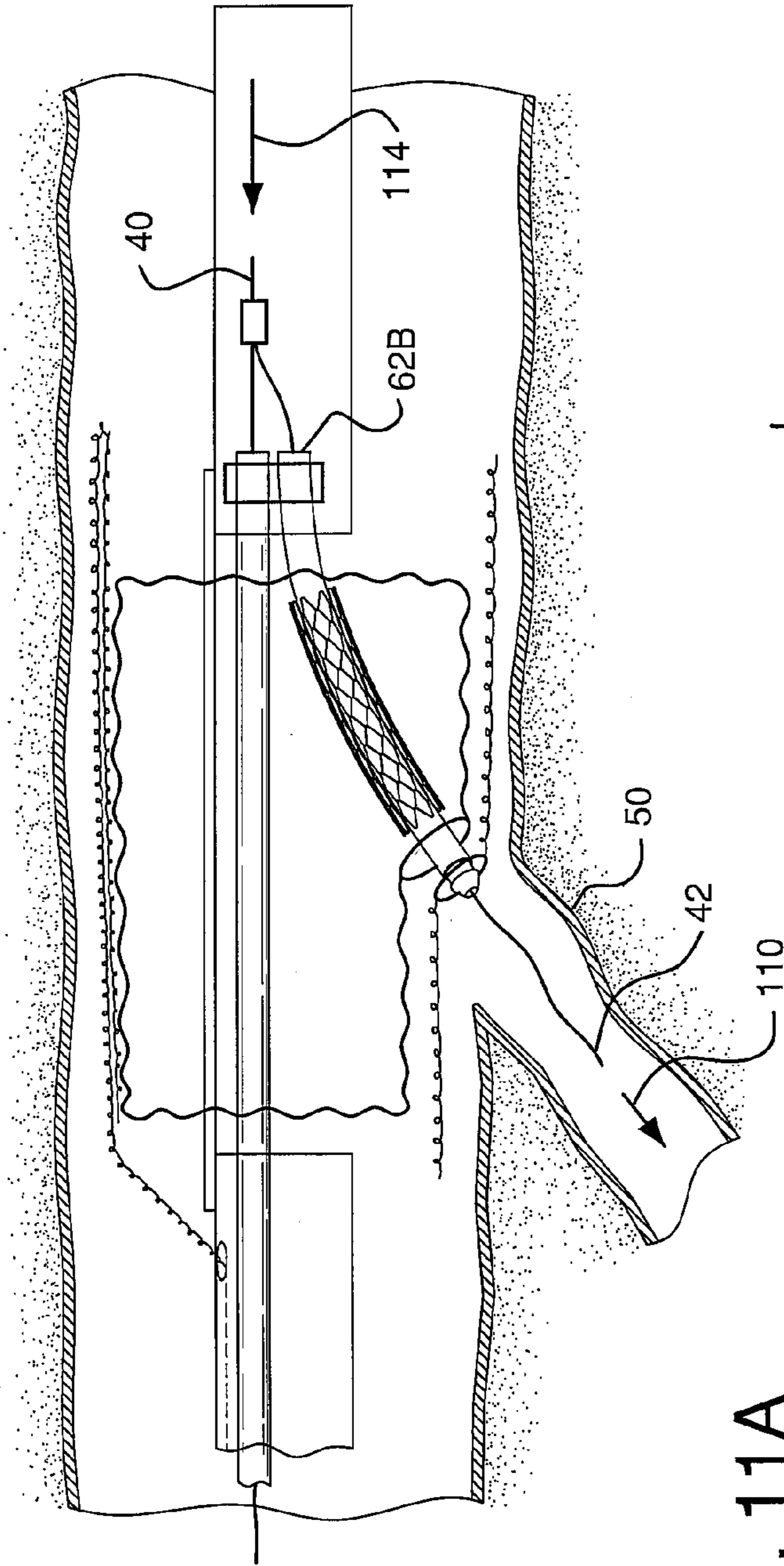


FIG. 11A

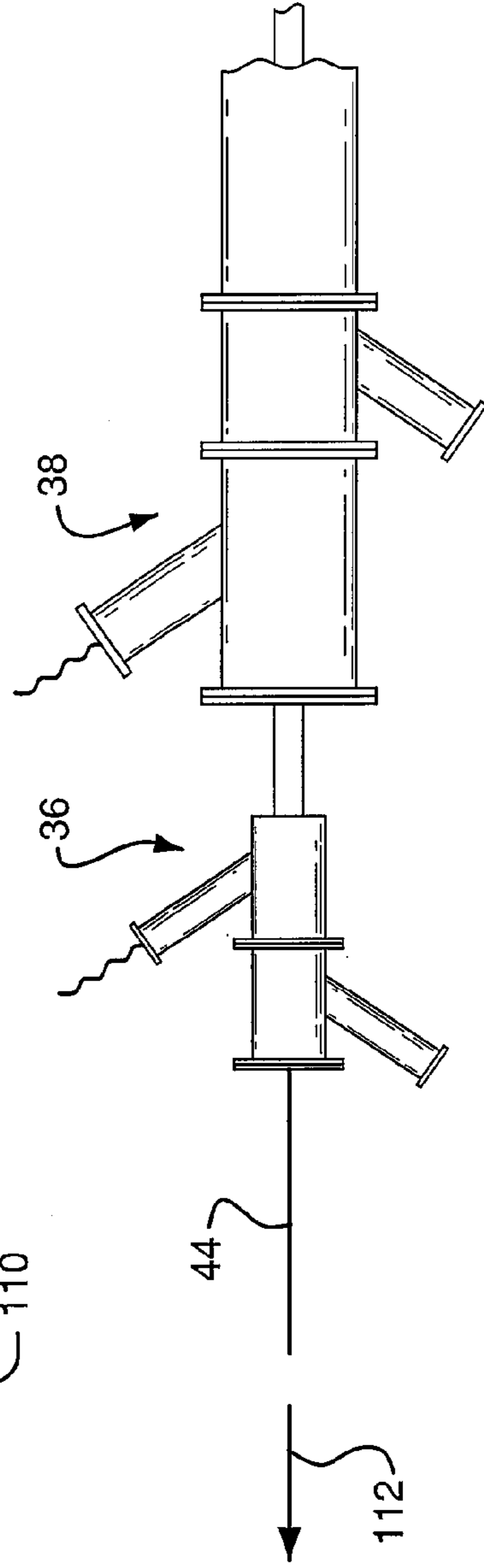


FIG. 11B

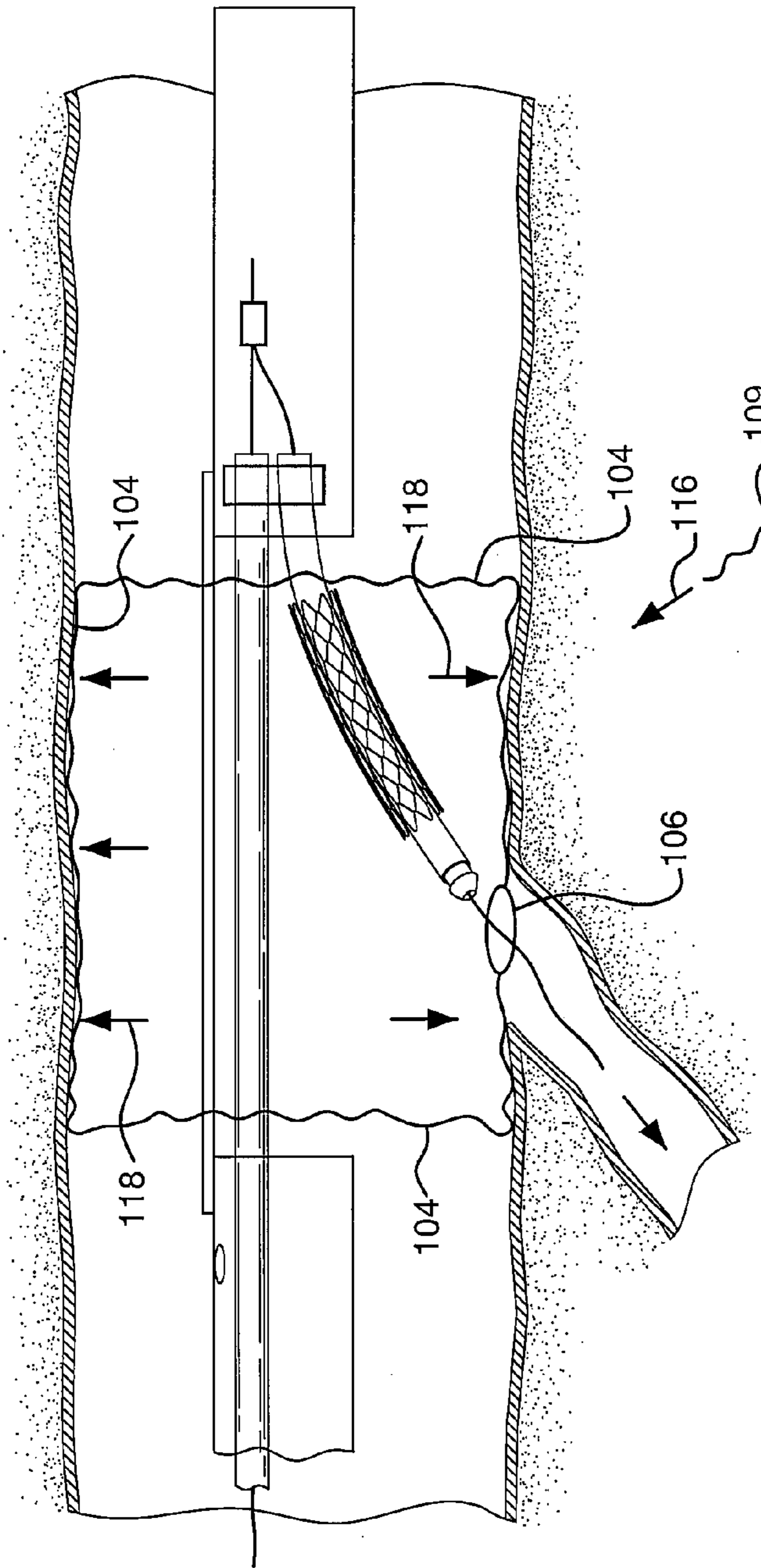


FIG. 12A

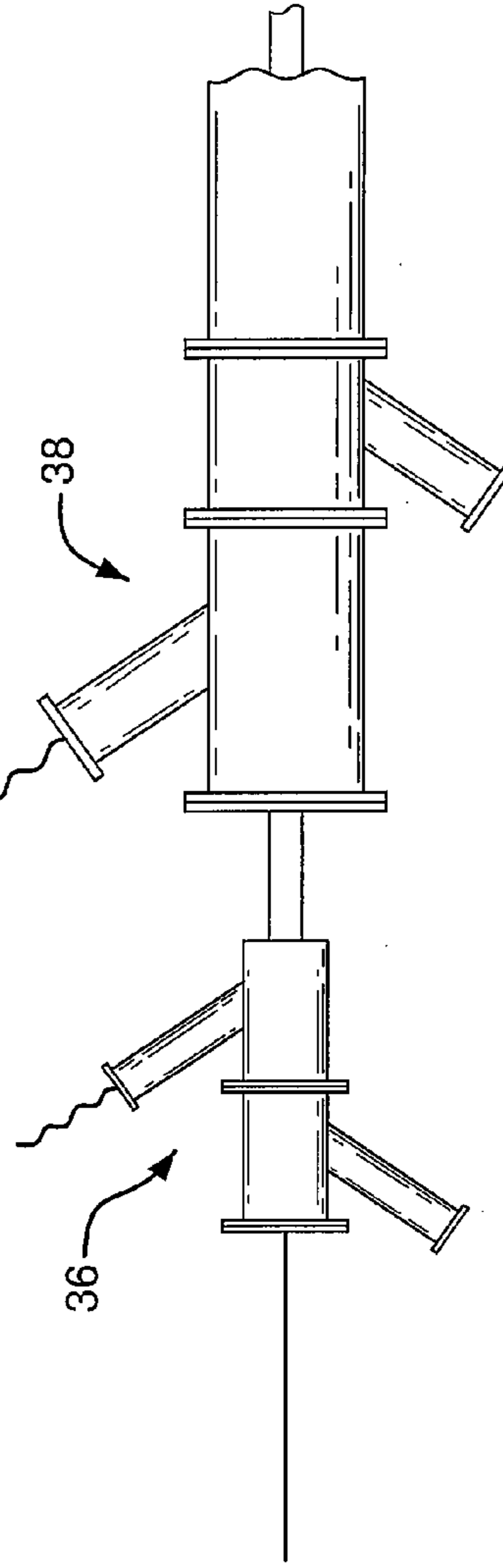


FIG. 12B

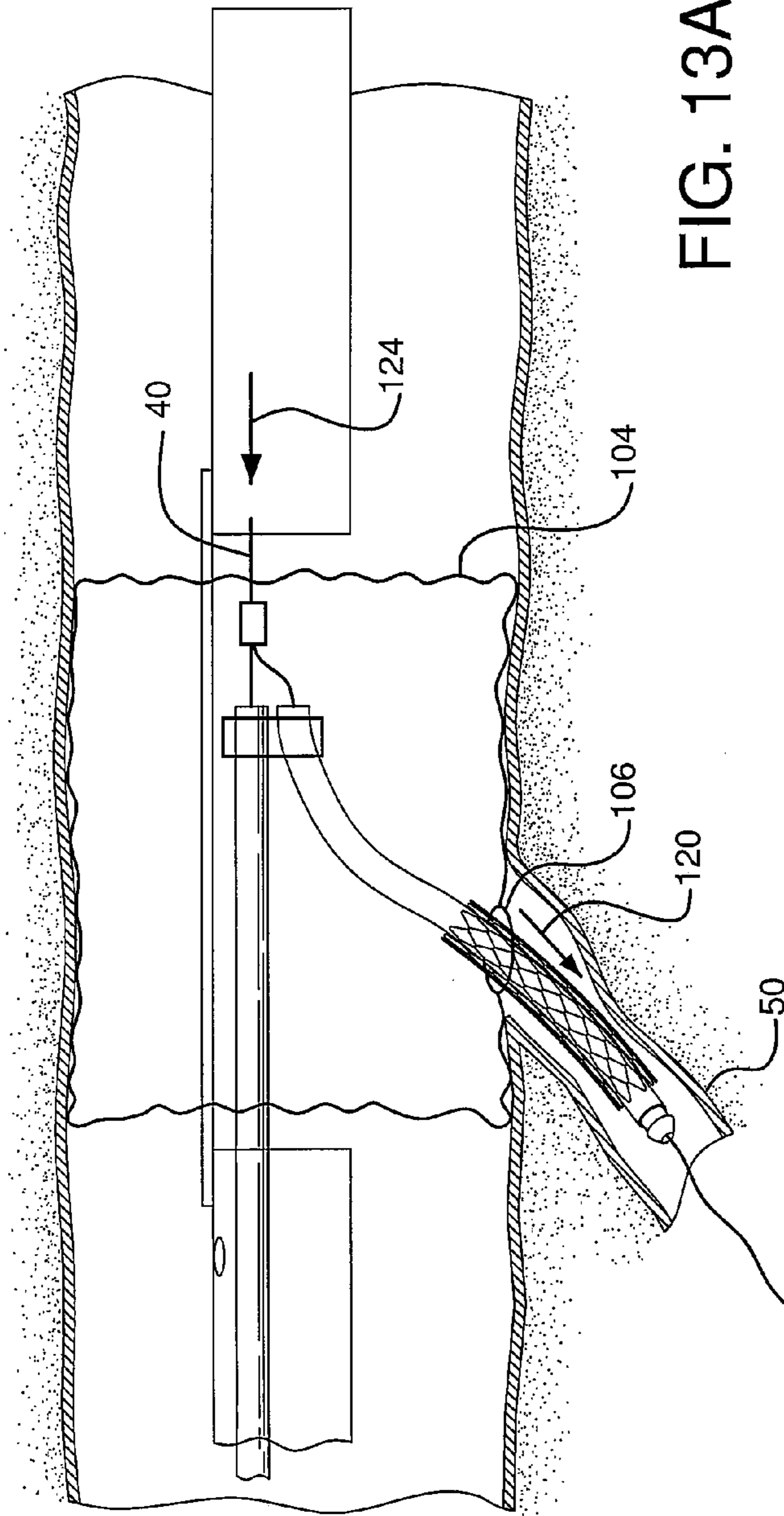


FIG. 13A

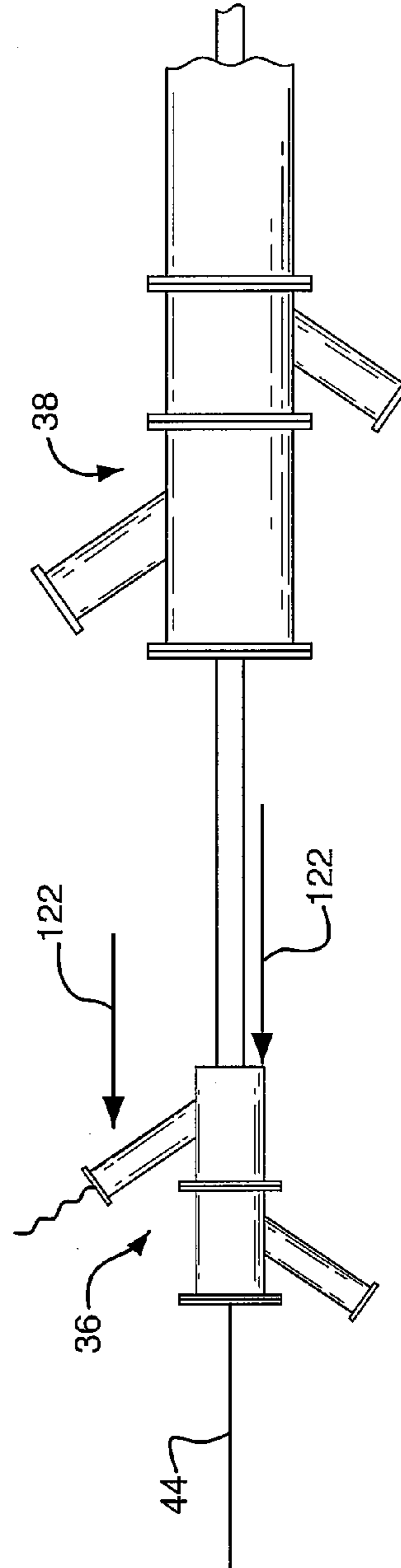


FIG. 13B

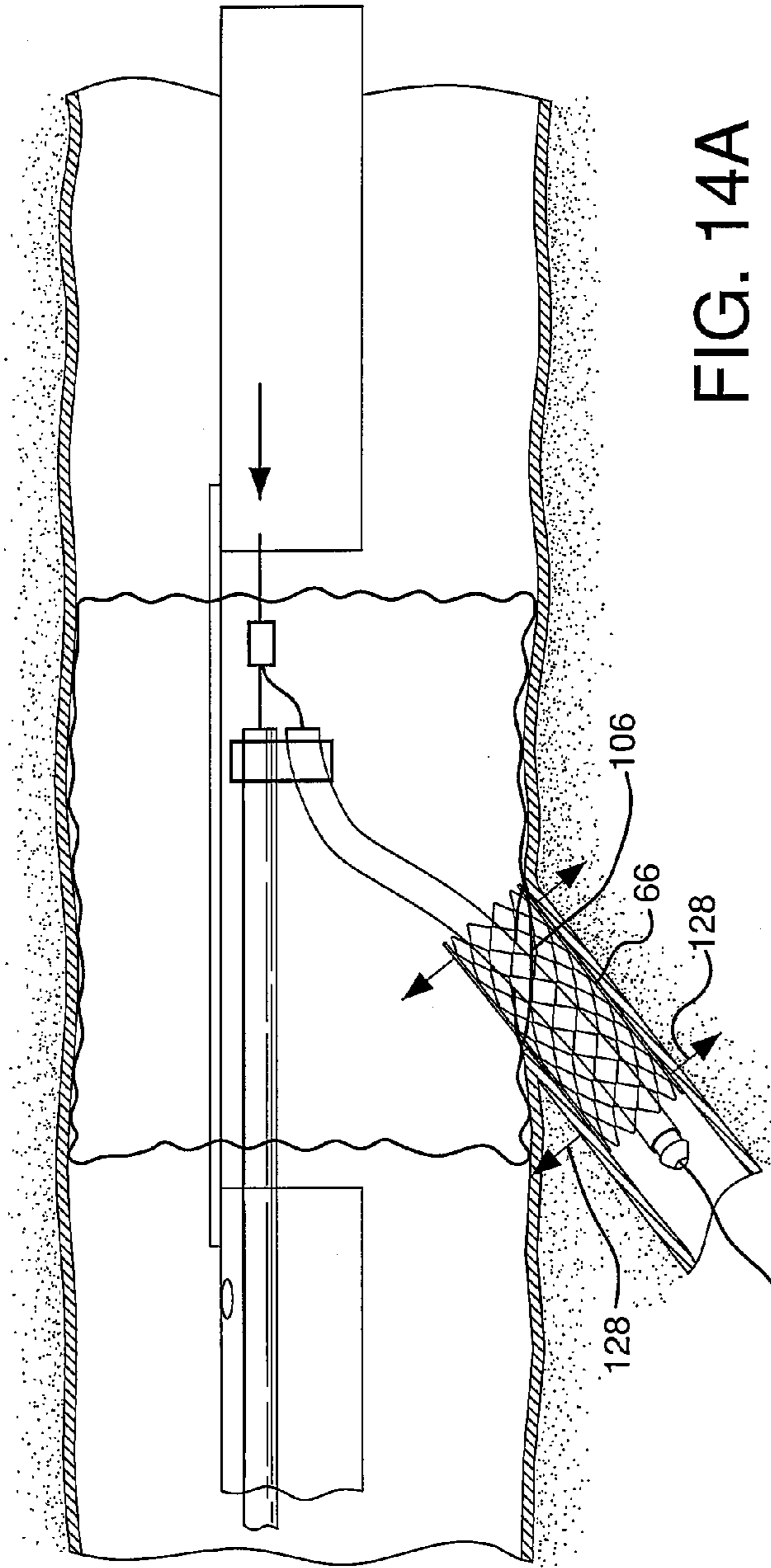


FIG. 14A

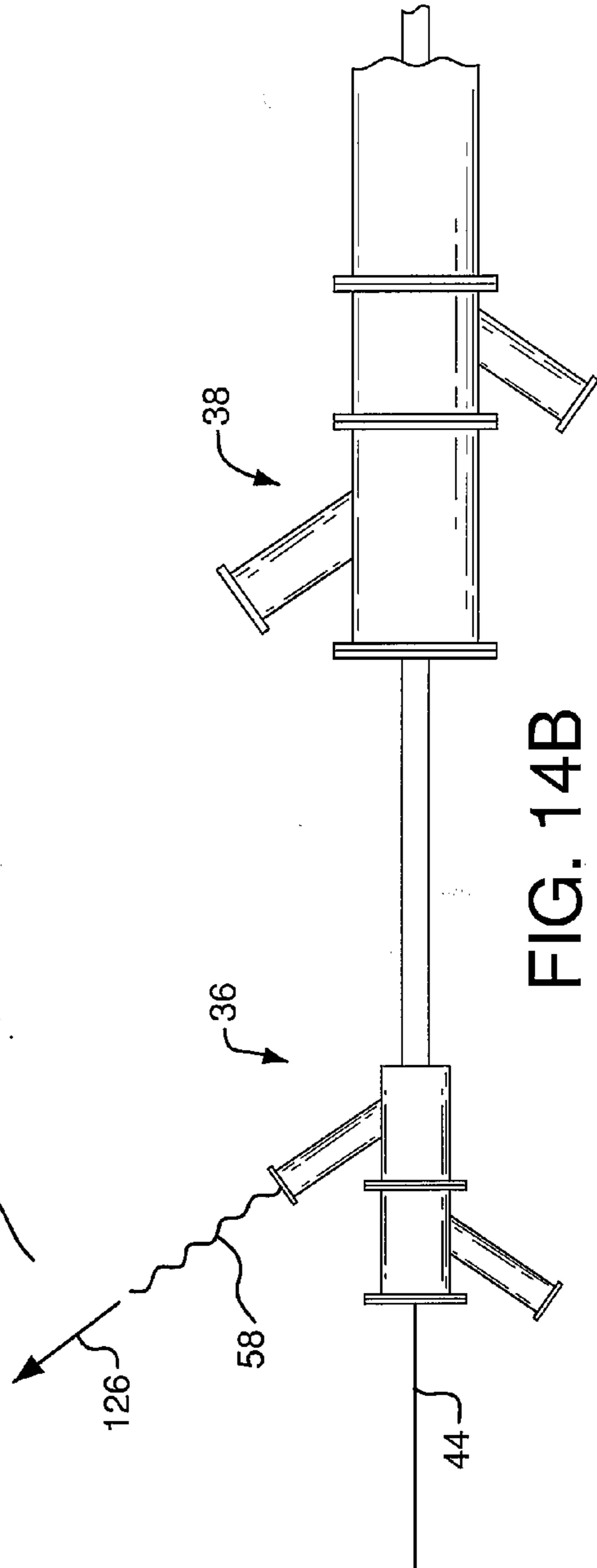


FIG. 14B

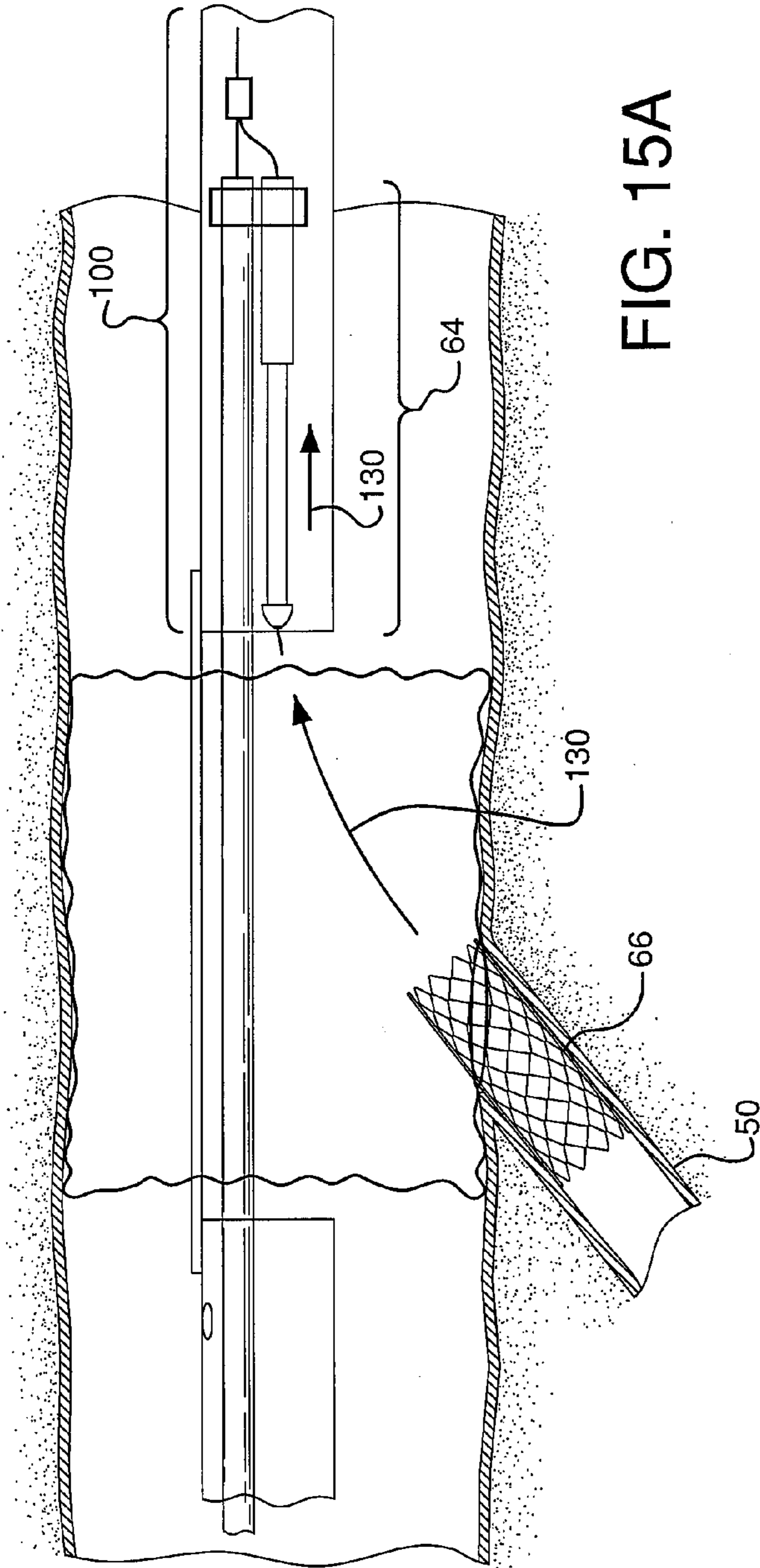


FIG. 15A

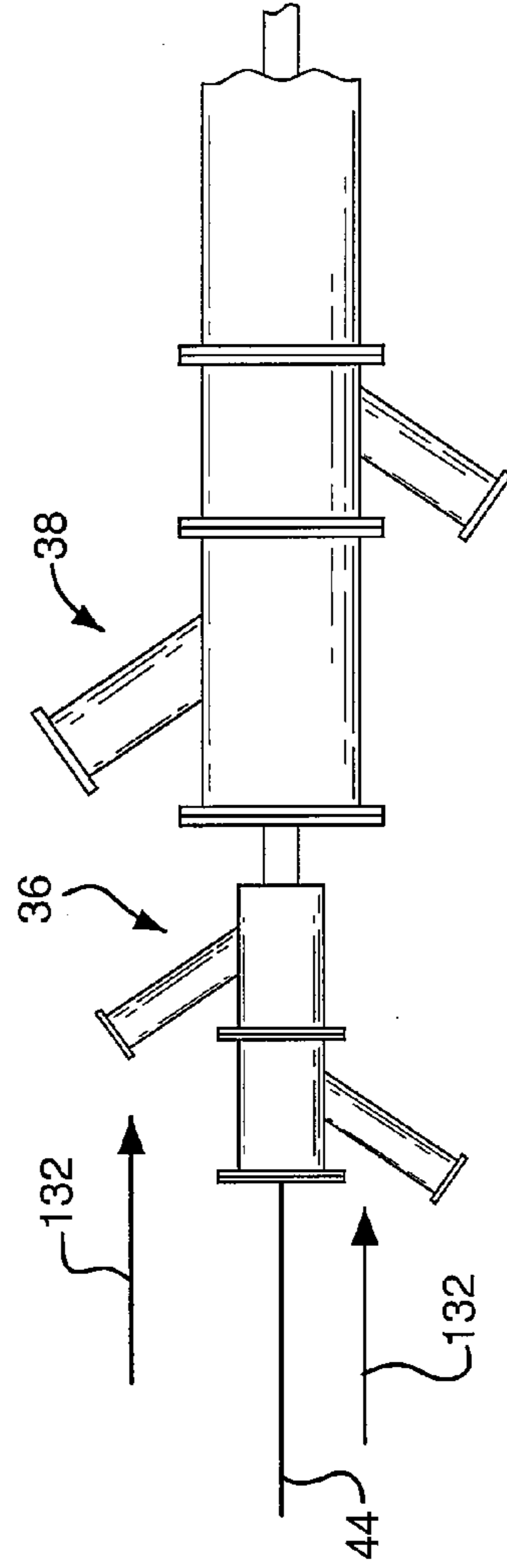


FIG. 15B

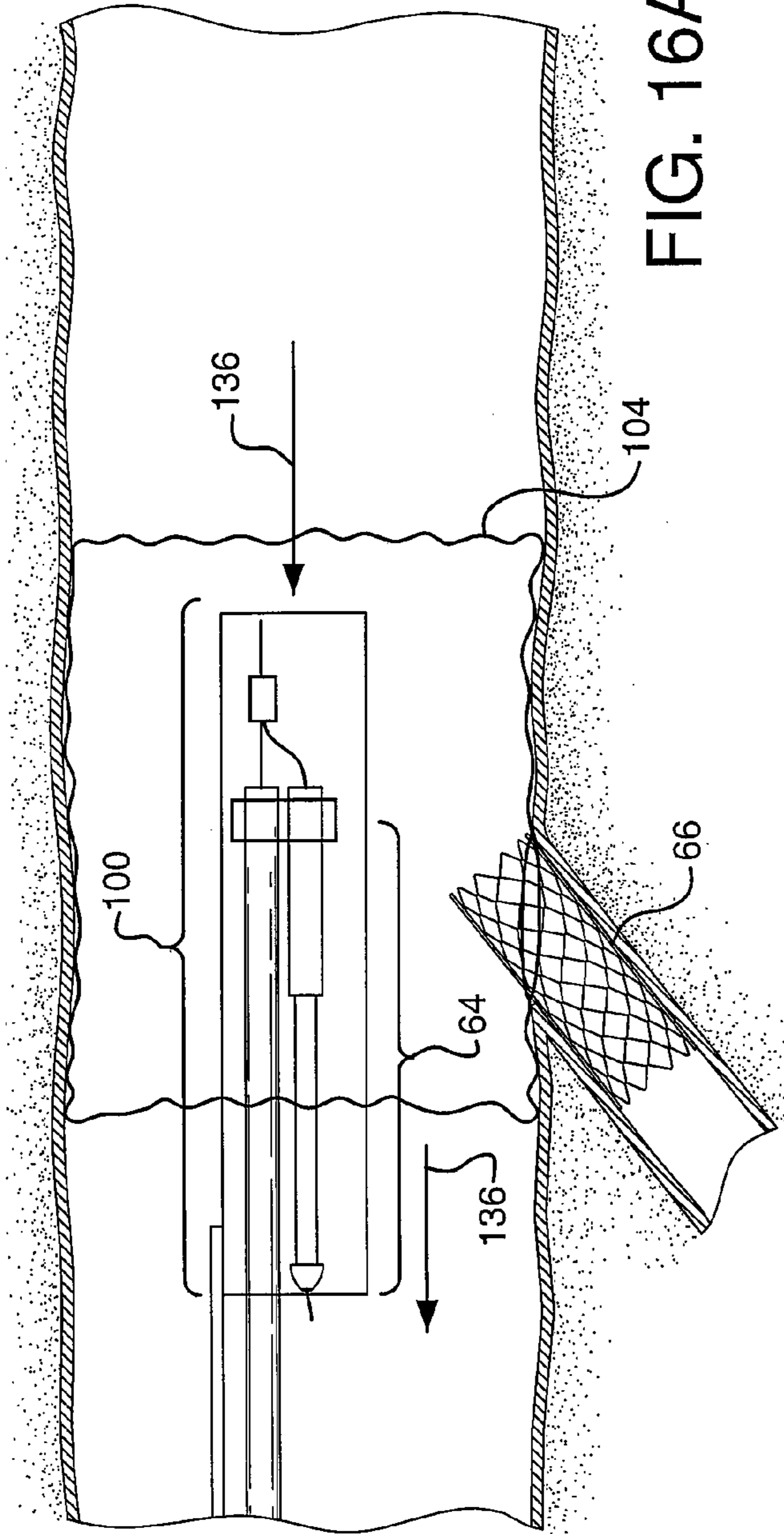


FIG. 16A

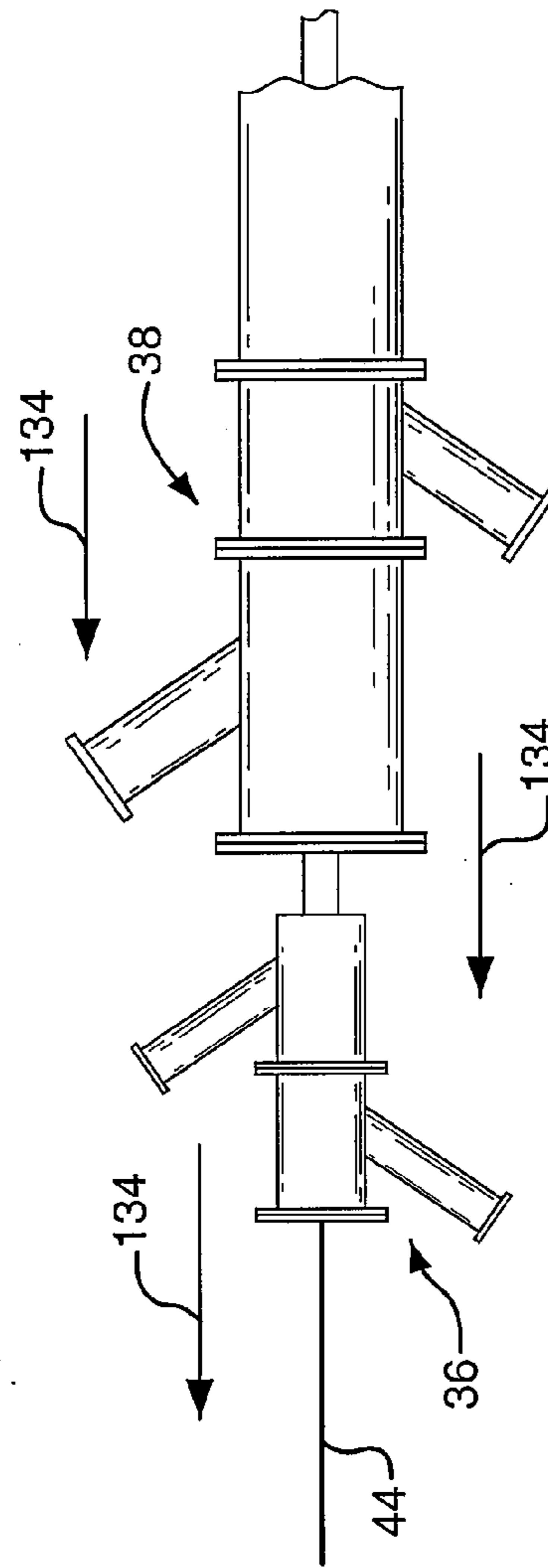


FIG. 16B

BRANCHED STENT DELIVERY SYSTEM

BACKGROUND OF THE INVENTION

The present invention generally relates to a delivery system and method for delivering an expandable endoluminal prosthetic device such as a stent graft and more particularly to a device and method for placing an acutely angled bifurcated stent graft through a single access incision. Expandable surgical devices such as stents or stent grafts are used in a variety of places in the human body to repair aneurysms and to support various anatomical lumens, such as blood vessels, respiratory ducts, gastrointestinal ducts, and the like.

Conventionally, these devices are deployed across an aneurysm or in the regions of a stenosis in the target body lumen to repair the aneurysm or to hold the lumen open. Because stent graft implantation is a relatively non-invasive procedure, it has been proven to be a favorable alternative to surgery in, for example, the repair of an aneurysm. Bifurcated devices with their trunk and branching configuration are particularly well suited for use in branching body lumen systems, such as in the coronary vasculature, and the peripheral vasculature. The coronary vasculature includes the right, left common, left anterior descending and circumflex arteries and their branches. The peripheral vasculature includes branches of the carotids, aorta, femoral, popliteal, internal iliac, or hypogastric and related arteries. Placement of such a bifurcated device can be rather complicated, and often involves approaching the bifurcated section of the artery through at least two side branches or through the trunk plus one side branch. The procedure is not only time consuming, but can also lead to more incision sites in the patient's body and can necessitate more complicated maneuvers for the surgeon. These complications are further exaggerated when an acutely angled or reverse direction side branch is accessed, as for example a repair of the hypogastric artery. U.S. Pat. No. 6,645,242 teaches a bifurcated intravascular stent graft comprising primary stent segments and a primary graft sleeve forming a main fluid channel and having a side opening therethrough.

However, there exists a need for a stent graft delivery system which would allow placement of bifurcated stent grafts into acutely angled vasculature such that simpler surgical procedures are enabled. A simplified surgical procedure would decrease the number or size of incisions, reduce the required surgical steps, and thereby reduce patient trauma associated with a more complex medical procedure.

SUMMARY OF THE INVENTION

The present invention further provides an interventional delivery system comprising: a first catheter having at its distal end a side branch vessel segment; a second catheter attached around the first catheter and having at its distal end a main vessel segment; and a side branch vessel device attached to the side branch vessel segment of the first catheter wherein the main vessel segment and the side branch vessel device are simultaneously delivered to a treatment site, and further wherein the second catheter has an opening in a side wall near the distal end of the second catheter to allow for passage of the side branch vessel segment of the first catheter. The second catheter may comprise a capture tube which surrounds the bifurcated guidewire and facilitates for the ease of bifurcated guidewire removal from a vessel. A bifurcated guidewire with at least two distal tips may be used with the first catheter. The two distal tips face opposing directions, wherein one of the two distal tips is the leading end and one of the tips is a reverse facing tip end.

The present invention further provides a first catheter having at its distal end a side branch vessel segment; a second catheter attached around the first catheter and having at its distal end a main vessel segment; a side branch vessel device attached to the side branch vessel segment of the first catheter; and a main vessel device attached to the main vessel segment of the second catheter. The main vessel device and the side branch vessel device are simultaneously delivered to a treatment site.

A method of deploying a branched stent assembly is also provided comprising: advancing a catheter assembly on a bifurcated guidewire to a treatment site; orienting the catheter assembly in the main vessel; pulling the bifurcated guidewire to orient the guidewire reverse facing tip into the side branch vessel; deploying the main vessel device in the main vessel; then advancing the side branch vessel device to a desired location; and deploying the side branch device. After stent deployment, removal of the delivery assembly is facilitated by advancing the guidewire and first catheter forward until the guidewire reverse facing tip and reverse facing portion of the first catheter are retracted from the side branch vessel allowing removal of the bifurcated guidewire along with the first and second catheters.

DESCRIPTION OF THE DRAWINGS

FIG. 1 shows the interventional delivery system comprising a first catheter and a second catheter upon insertion in a vessel.

FIGS. 2A and 2B show the first catheter of the interventional delivery system. FIG. 2A depicts a bent shaft configuration and FIG. 2B depicts a shaft configuration using a connector.

FIG. 3 shows the bifurcated guidewire assembly with a leading segment and a reverse facing segment.

FIG. 4A shows a first catheter with a bent shaft and apex opening for the bifurcated guidewire with a side branch device mounted on the side branch vessel segment of the first catheter shaft.

FIG. 4B shows a first catheter with a shaft configuration using a connector for the bifurcated guidewire with a side branch device mounted on the side branch vessel segment of the first catheter shaft.

FIG. 5 shows an enlarged view of the side branch vessel segment with a side branch vessel device mounted and constrained within a sheath.

FIGS. 6A and 6B show side views of a second catheter with a side branch opening.

FIG. 7 is an isometric view of an expanded main body stent graft.

FIG. 8 is a partial cross-sectional view of a main body stent with a bifurcated guidewire and a first catheter with a side branch device.

FIG. 9 is a partial cross-sectional view of a main body stent with a bifurcated guidewire and a first catheter with a side branch device contained in a second catheter. Also shown is a constraint sheath over the main body stent and the apertures in the main body stent and the constraining sheath.

FIGS. 10A and 10B show partial cross-sectional views of the distal device portion and the proximal hub portions of the interventional delivery system of the present invention. The distal device portion is positioned within a main vessel adjacent to a branched vessel.

FIGS. 11A and 11B show partial cross-sectional views of the distal device portion and the proximal hub portions of the interventional delivery system of the present invention. The

reverse facing guidewire is shown being advanced into the side branch, acutely angled vessel.

FIGS. 12A and 12B show partial cross-sectional views of the distal device portion and the proximal hub portions of the interventional delivery system of the present invention. The main body stent is shown in an expanded state.

FIGS. 13A and 13B show partial cross-sectional views of the distal device portion and the proximal hub portions of the interventional delivery system of the present invention. The reverse facing segment of the first catheter with a constrained side branch device is shown being advanced into the side branch, acutely angled vessel.

FIGS. 14A and 14B show partial cross-sectional views of the distal device portion and the proximal hub portions of the interventional delivery system of the present invention. The side branch stent is shown in an expanded state.

FIGS. 15A and 15B show partial cross-sectional views of the distal device portion and the proximal hub portions of the interventional delivery system of the present invention. The reverse facing portion of the first catheter and the guidewire are shown being advanced into a capture tube.

FIGS. 16A and 16B show partial cross-sectional views of the distal device portion and the proximal hub portions of the interventional delivery system of the present invention. The first catheter, the second catheter, and the guidewire are shown being withdrawn from the treatment site.

DETAILED DESCRIPTION OF THE INVENTION

The present invention provides an interventional delivery system for the placement of bifurcated stent grafts into acutely angled vasculature. Acutely angled vasculature may exist in renal vessels, subclavian arteries, biliary ducts, prostate vessels, and other non-vascular applications as well. The challenge in stent placement is deployment from a main vessel such as a femoral artery to a reverse acute angle vessel. The present invention provides a device and procedure which decreases the number and size of incisions required to place bifurcated stent grafts into acutely angled vasculature, and further reduces the required surgical steps and patient trauma associated with this traditionally more complex medical procedure. As shown in FIG. 1, the present invention provides an interventional delivery system 30 comprising a first catheter shaft 32, a second catheter assembly 34A or 34B, a first catheter hub assembly 36, a second catheter hub assembly 38, a bifurcated guidewire leading tip 40, a bifurcated guidewire reverse facing tip 42, a bifurcated guidewire proximal tip 44, and a device assembly 46. The interventional delivery system 30 is shown positioned in an anatomical main vessel 48 so that the device assembly 46 is positioned approximate to an anatomical side branch vessel 50. As described in subsequent figures, the device assembly 46 will be deployed to form a main body stent within the main vessel 48 along with an integrated side branch stent within the side branch vessel 50.

Shown in FIGS. 2 through 9 are various sub-components and assemblies of the interventional delivery system 30 (of FIG. 1). Shown in FIG. 2A is a first catheter assembly 52A having a first catheter hub assembly 36. The hub assembly 36 includes a perfusion port 54, a bifurcated guidewire port 56, a side branch deployment line 58 protruding from a deployment line port 60. The first catheter assembly 52A further comprises a first catheter shaft 32 that has an apex opening 62A. Apex opening 62A as shown is a cut-opening through the wall of the first catheter shaft 32. The first catheter shaft is shown bent about the apex opening 62A, forming a reverse

facing segment 64. The reverse facing segment 64 has a side branch device portion 66 and a side branch device to apex opening separation length 68.

As shown in FIG. 2B a first catheter assembly 52B has a first catheter hub assembly 36. The hub assembly 36 includes a perfusion port 54, a bifurcated guidewire port 56, a side branch deployment line 58 protruding from a deployment line port 60. The first catheter assembly 52B further comprises a first catheter shaft 32 that has an apex opening 62B. Apex opening 62B as shown comprises the open ends of a cut catheter shaft 32. The two cut ends are joined at connection 70. The two cut shafts as shown form a reverse facing segment 64. The reverse facing segment 64 has a side branch device portion 66 and a side branch device to apex opening separation length 68.

Depicted in FIG. 3 is a bifurcated guidewire assembly 72 having a proximal tip 74 and a main segment 76. Within the distal portion of the guidewire main segment 76 is a connection 78, defining a leading guidewire segment 80 and a reverse facing guidewire segment 82. The leading guidewire segment has a leading tip 84 and the reverse facing guidewire segment has a reverse facing tip 86.

FIG. 4A shows a first catheter assembly (52A of FIG. 2A) combined with a bifurcated guidewire assembly (72 of FIG. 3). Referring to FIGS. 2A, 3, and 4A, shown is a bifurcated guidewire assembly 72 positioned within a first catheter assembly 52A. Shown protruding from an apex opening 62A is the bifurcated guidewire connection 78 along with the leading guidewire segment 80. The proximal end 74 of the bifurcated guidewire protrudes from the bifurcated guidewire port 56 and the reverse facing tip 86 of the guidewire protrudes from the reverse facing segment of the first catheter.

Similarly, FIG. 4B shows a preferred first catheter assembly (52B of FIG. 2B) combined with a bifurcated guidewire assembly (72 of FIG. 3). Referring to FIGS. 2B, 3, and 4B, shown is a bifurcated guidewire assembly 72 positioned within a first catheter assembly 52B. Shown protruding from an apex opening 62B is the bifurcated guidewire connection 78 along with the leading guidewire segment 80. The proximal end 74 of the bifurcated guidewire protrudes from the bifurcated guidewire port 56, and the reverse facing tip 86 of the guidewire protrudes from the reverse facing segment of the first catheter. The tube-to-tube connection 70 can include a friction-reducing component or feature to allow the deployment line 58 to easily slide against the tubes or apex opening as the deployment line is activated.

Shown in FIG. 5 are a partial cross-sectional view of the reverse facing segment 64 that includes a side branch device portion 66 and a side branch device to apex opening separation length 68. Shown is the bifurcated guidewire 72 reverse facing tip 44 exiting from an olive 88. Positioned onto a side branch accommodating segment 94 is a constrained, self-expanding side branch device 90. The side branch device 90 is held in a compressed state by a constraining sheath 92. Attached or integral to the constraining sheath is a side branch device deployment line 58.

FIGS. 6A and 6B are side views of two embodiments of a second catheter. Shown in FIG. 6A is a second catheter assembly 34A having a second catheter hub assembly 38. The second catheter hub assembly further includes a proximal perfusion port 54. The hub assembly is joined to a second catheter main body 96. Near the distal end of the second catheter main body 96 is a side branch device opening 98, formed by a cut-out portion of the catheter wall. The opening 98 allows a bifurcated guidewire and a side branch device to be subsequently advanced from the second catheter. After deployment, the bifurcated guidewire can be pulled through

5

the opening **98** into the second catheter for removal. At the distal end of the second catheter main body is a capture tube portion **100**. This tube portion “captures” the bifurcated guidewire after device deployment, allowing for a non-traumatic removal of the guidewire and delivery system.

Similarly, FIG. **6B** depicts an alternate embodiment of a second catheter assembly **34B**. The distal end of the second catheter main body **96** is joined to the capture tube portion **100** by at least one main body to capture tube joining member **102**. The main body **96** and the capture tube **102** are therefore separated and connected by the joining members **102**. The gap between the main body and the capture tube forms an opening **98** functionally similar to the opening **98** shown in FIG. **6A**.

FIG. **7** is an isometric view of an expanded main body device **104**. An aperture **106** is formed in the main body device wall, permitting a side branch device to be subsequently inserted through and attached to the aperture/main body.

FIG. **8** is a partial cross-sectional view of a main body device **104** surrounding a first catheter assembly **52B**. A bifurcated guidewire **72** is positioned within the first catheter (as previously shown in FIG. **4B**). A reverse facing portion of the first catheter having a constrained side branch device is shown protruding through an aperture **106** in the main body stent. Exiting from the reverse facing portion of the first catheter is the reverse facing tip **42** of the bifurcated guidewire. Also shown are the first catheter shaft **32** and the apex opening **62B**.

FIG. **9** is a partial cross-sectional view of the components depicted in previous FIG. **8** along with a second catheter **34B** (refer to FIG. **6B**). Shown is a second catheter main body **96**, connected to a capture tube portion **100** by at least one joining member **102**. The distal end of the bifurcated guidewire is shown positioned within the capture tube portion **100**. The first catheter shaft **32** is shown positioned within the second catheter main body **96**. Also shown are a constraining sheath **92** and the attached or integral main body deployment line **109**. The reverse facing portion of the first catheter is shown protruding through an aperture **108** within the constraining sheath **92**.

A sequence used to deliver and deploy main body and side branch stents according to the present invention is depicted in FIGS. **10** through **16**.

Deployment Step 1

FIG. **10A** is a partial cross-sectional view of the distal end of an interventional delivery system similar to that of FIG. **1**. A device assembly is shown initially positioned in an anatomical main vessel **48** so that the device assembly is positioned approximate to an anatomical side branch vessel **50**. Shown are a bifurcated guidewire leading tip **40** and a bifurcated guidewire reverse facing tip **42**. The device assembly (**46** of FIG. **1**) has been expanded to display the internal components as shown in FIG. **9**. FIG. **10B** depicts the proximal end of the interventional delivery system, similar to that shown in FIG. **1**. Shown are a first catheter hub assembly **36** and a second catheter hub assembly **38**.

Deployment Step 2

FIGS. **11A** and **11B** show the bifurcated guidewire reverse facing tip **44** being advanced into the side branch vessel **50** along the direction indicated by arrow **110**. The guidewire reverse facing tip **42** is advanced by pulling (in direction indicated by arrow **112**) on the proximal end **44** of the guidewire. The two hub assemblies **36**, **38** are held stationary as the proximal end of the guidewire is pulled. As the proximal end of the guidewire is pulled, the guidewire leading tip

6

40 is advanced towards the apex opening **62B** in the direction shown by arrow **114**. The guidewire reverse facing tip **42** is therefore forced to advance into the side branch vessel **50** in the direction of arrow **110**.

Deployment Step 3

Referring to FIGS. **12A** and **12B**, the main body stent **104** is deployed by pulling on the main body stent deployment line **109** in the direction indicated by arrow **116**. By releasing the constraining sheath (**92** of FIG. **9**) the main body stent is allowed to self-expand in the directions indicated by arrows **118**. The two hub assemblies **36**, **38** are held stationary as the deployment line is pulled. Note that the guidewire and/or the side branch device are positioned through the aperture **106** in the main body device **104**.

Deployment Step 4

The side branch device is then advanced into the side branch vessel, as depicted in FIGS. **13A** and **13B**. The side branch device is advanced along the direction indicated by arrow **120** by holding stationary the second catheter hub assembly **38** while concurrently pulling on the guidewire **44** and the first catheter hub assembly **36**. The guidewire may be optionally locked onto the first catheter hub assembly **36** to facilitate this step. As the guidewire and hub assembly are pulled, the distal tip of the guidewire **40** is pulled in the direction indicated by arrow **124**, forcing the side branch device to advance partially through the main body device aperture **106** and into the side branch vasculature **50** in the direction **120**.

Deployment Step 5

As shown in FIGS. **14A** and **14B**, the side branch deployment line **58** is then pulled in the direction indicated by arrow **126**, allowing the side branch device **66** to self-expand as indicated by arrows **128**. Note that the side branch device is partially contained within and constrained by the main body device aperture **106**. The two hub assemblies **36**, **38** are held stationary as the deployment line is pulled.

Deployment Step 6

Referring to FIGS. **15A** and **15B**, the delivery system of the present invention is withdrawn from the vasculature by forcing the reverse facing portion of the first catheter **64** out of the expanded side branch device and into the capture tube **100** along the direction as indicated by arrows **130**. The first catheter reverse facing portion is driven into the capture tube by pushing the first catheter hub assembly **36** along with the guidewire **44** along the direction as shown by arrows **132**. The second catheter hub assembly **38** is held stationary as the first catheter hub assembly and the guidewire are advanced.

Deployment Step 7

To complete the delivery of the devices and systems of the present invention, the first catheter hub assembly **36**, the guidewire proximal tip **44**, and the second catheter hub assembly **38** are concurrently pulled in the direction as shown by arrows **134** of FIGS. **16A** and **16B**. The capture tube **100**, containing the bifurcated guidewire and the reverse facing portion or the first catheter **64** are non-traumatically removed from the vasculature, leaving the expanded main body device **104** and the attached side branch device **66** in the vasculature.

Referring back to FIG. **7**, the main body device **104** is shown with a single side-wall aperture **106**. In an alternate configuration, a main body device can have two, three, four, five, six or more side branch apertures. The various catheters of the present invention can incorporate more than one device; for example, a first catheter can incorporate two or more side branch devices. The sealing or interference fit

between a main body and a side branch device can be enhanced by the incorporation of a “sealing sleeve”. See for example U.S. Pat. No. 6,645,242 to Quinn for a disclosure of such sealing sleeves. Multiple sealing sleeves can be incorporated into a main body device to enhance the sealing or attachment of multiple side branch devices. Sealing sleeves can be “internal to” or “external to” the lumen of a main body stent and can be shaped and sized to seal a specifically configured side branch device.

Stents used in the present invention can be bare (uncovered), coated with a variety of drug eluting, anti-thrombogenic or other coatings, or can include a partial or full cover (as in a stent graft). Anchoring mechanisms, such as barbs, “fish-scales”, biological attachment means, or other features can be incorporated into the main body and/or a side branch device to facilitate anchoring to the vasculature.

Main body stents and/or side branch stents can have a uniform profile or have non-uniform profiles such as tapers, “trumpet-end” shapes, “dog-bone” shapes, curves or other profiles that enhance the device performance within a particular treatment site. Multiple devices of the present invention can be “ganged” or interconnected to form a multi-component system. Devices of the present invention can include features that allow or enhance the interconnection or “docking” between multiple devices.

Radiopaque markers or indicators can be incorporated into a main body device, the various catheters used in the present invention and/or a side branch device to facilitate placement and visualization within the vasculature.

Devices of the present invention can be used to treat non-vascular conduits, hollow or tubular parts of organs, such as biliary, bladder, urethra, gastrological, bronchi, bile, and other ducts. Devices of the present invention are particularly suited for, but not limited to, side branch vessels that have an “acute” angle from the main body (see for example FIG. 1).

Devices of the present invention can be balloon-expandable as well as self-expanding. For example, the first catheter according to the present invention can incorporate a balloon (or balloons) and inflation lumens as required to expand a particular device. Combinations of self-expanding and balloon-expandable devices can be configured according to the present invention. Also, separate balloon expanders can be used within the scope of the present invention.

Catheter components of the present invention can be fabricated from common materials such as nylons, polycarbonates, polyethylenes, polypropylenes, polytetrafluoroethylenes, polyvinyl chlorides, polyurethanes, polysiloxanes, stainless steels, nitinol, or other biocompatible materials.

While particular embodiments of the present invention have been illustrated and described herein, the present invention should not be limited to such illustrations and descriptions. It should be apparent that changes and modifications

may be incorporated and embodied as part of the present invention within the scope of the following claims.

The invention claimed is:

1. An interventional delivery system for acutely angled branched stent grafts comprising: a first catheter having at its distal end a side branch vessel segment; a second catheter attached around the first catheter and having at its distal end a main vessel segment; a side branch vessel device attached to the side branch vessel segment of the first catheter; a main vessel device attached to the main vessel segment of the second catheter; and a bifurcated guidewire with at least two distal tips which is configured to move slidably within the first catheter, wherein the main vessel device and the side branch vessel device are simultaneously delivered to a treatment site; wherein the first catheter further comprises a crotch which provides a positive stop for maintaining a portion of the side branch device within a sealing sleeve of the main vessel device body to ensure adequate engagement.

2. A method of deploying a branched stent assembly comprising:

advancing a catheter assembly comprising: an interventional delivery system for acutely angled branched stent grafts comprising: a first catheter having at its distal end a side branch vessel segment; a second catheter attached around the first catheter and having at its distal end a main vessel segment; a side branch vessel device attached to the side branch vessel segment of the first catheter; a main vessel device attached to the main vessel segment of the second catheter; and a bifurcated guidewire with at least two distal tips which is configured to move slidably within the first catheter, wherein the main vessel device and the side branch vessel device are simultaneously delivered to a treatment site, on a bifurcated guidewire to a treatment site;

orienting the catheter assembly in the main vessel; pulling the bifurcated guidewire to orient a portion of the bifurcated guidewire into the side branch vessel; deploying the main vessel device in the main vessel; advancing the side branch vessel device to a desired location; deploying the side branch device; pushing the guidewire forward until the portion of the guidewire in the side branch vessel is retracted into the first catheter; pushing the first catheter forward until the reverse facing tip is retracted into the capture tube; and withdrawing the first catheter, the bifurcated guidewire, and the second catheter.

3. The method of claim 2 wherein the branched stent assembly is deployed in a main vessel and side branch vessel which are oriented at an acute angle to each other.

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